



Biological Agent Sensor Testbed*

MIT Lincoln Laboratory

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Biological Agent Sensor Testbed (BAST) Development Status



Biological Agent Sensor Testbed

- **Motivation and Goals**
- **Sensor Concept**
- **Breadboard Apparatus and Data**
- **Engineering Development Unit Design**
 - **Optical**
 - **Electrical**
 - **Mechanical**
 - **Algorithm**
- **Summary**



Motivation and Goals

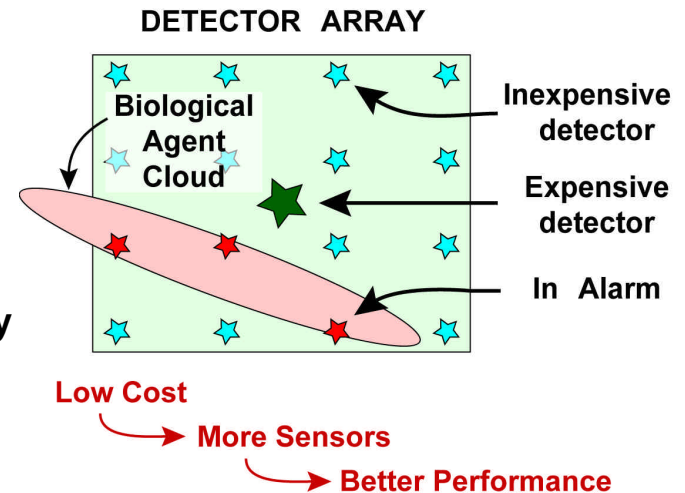
- The cost of current biological agent detection systems strongly limits the number of locations that can be protected.

- Needed: biological agent detection system that can be made in large quantities for the protection of many locations

- Low cost
- Good performance
sensitivity, false positive rate, probability of detection, response time
- Willing to trade performance for cost, size, weight, and power

- **BAST Goals**

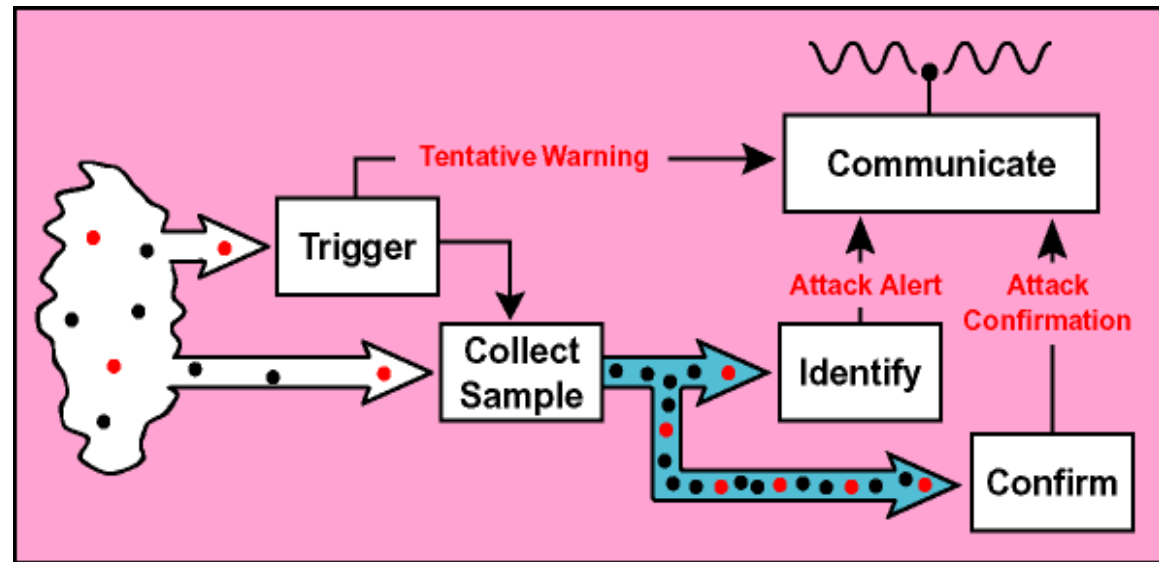
- Low manufacturing cost (<\$1000)
- Low false positive rate (<1/Week)
- Sensitivity < 500 ppl (98% detection confidence)
- Response time < 60 sec
- Small size (0.5 cu. ft.), low weight (<20 lbs)





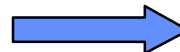
Biological Agent Detection

Typical Bio-sensor Architecture

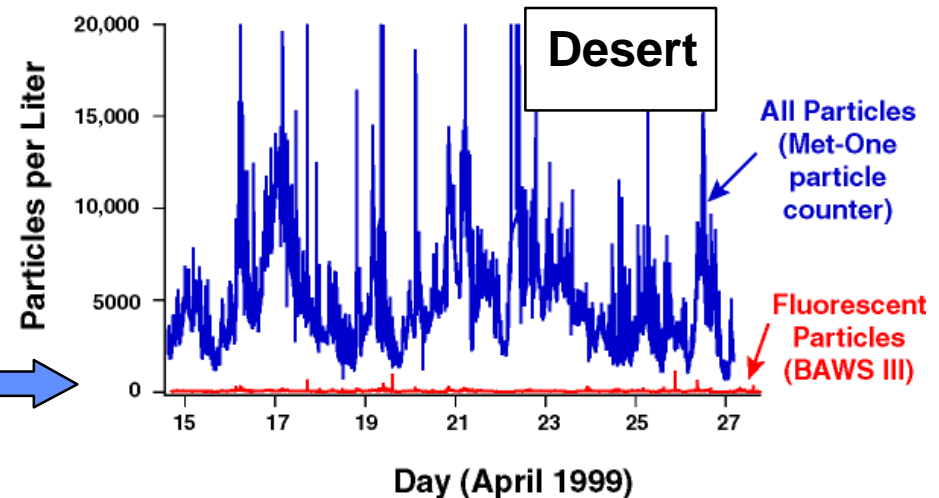


Fluorescence Trigger

- Sensitive
- High probability of detection
- Fast
- Relatively low false positive rate



Fluorescent vs. Total Particles





UV-LED-Based Biological Agent Detection

- **Advantages**

- **Cost**

- Enable low cost and small size sensors

- **Performance**

- Affordable large array networks for improved sensitivity

- Multiple operating wavelengths for improved discrimination

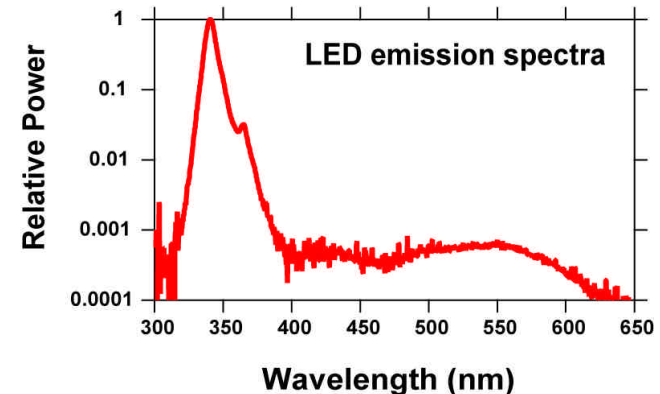
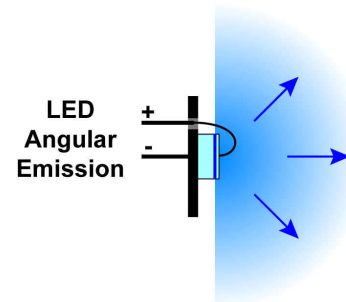
- **Issues**

- **Short wavelengths**

- **LED lifetime**

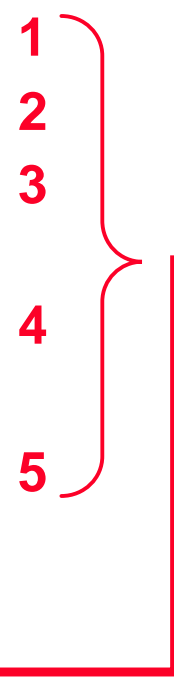
- **Low brightness**

- **Relatively broad spectral emission**



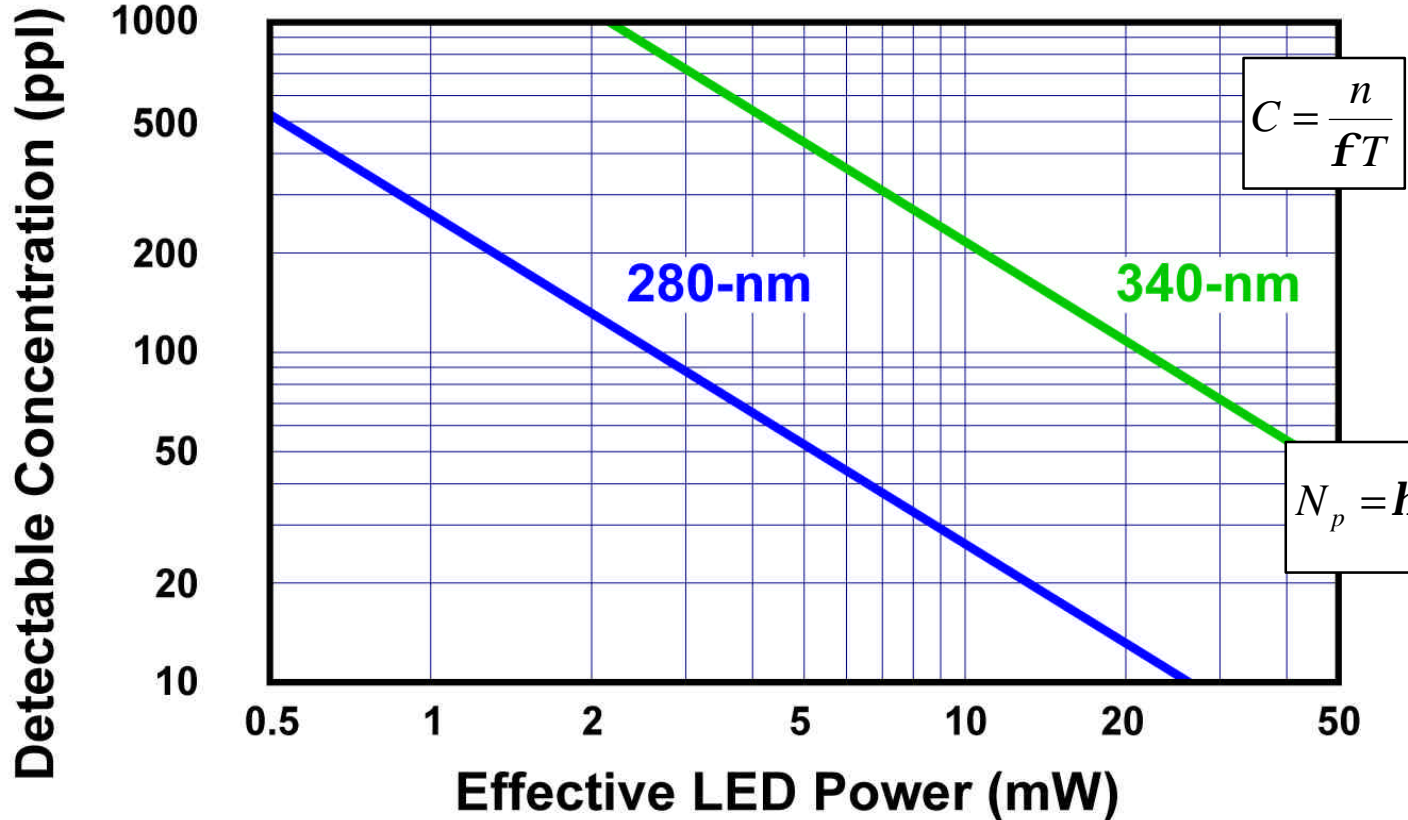


Sensor Concept

- Particle illumination with 280-nm and 340-nm LED radiation and with 820-nm laser diode radiation
 - Detection of 280-nm and 340-nm LED induced side fluorescence
 - 300 – 400 nm (280-nm excitation)
 - 400 – 500 nm (280-nm excitation)
 - 400 – 500 nm (340-nm excitation)
 - Detection of side elastic scattering (280 nm)
 - Detection of forward elastic scattering at 820-nm
 - Measure particle size
 - 5 independent measurement channels
- 
- A red bracket on the right side of the list groups the first five items (the three fluorescence detection items, the side elastic scattering item, and the forward elastic scattering item) together. A red arrow points from this bracket to the '5 independent measurement channels' item, indicating that these five items correspond to the five channels.



Fluorescence Detection Analysis



| | |
|---|--|
| Number of detected particles (N_b) | 100 particles |
| Particle fluorescence cross-section (s) | $(50, 5) \times 10^{-12} \text{ cm}^2$ @ (280, 340) nm |
| Number of detected photo-electrons (N_{pe}) | 100 photoelectrons |
| Photon collection, detection efficiency (h_d) | 30% , 15% |
| Threat detection time (T) | 60 s |



LED Sources

280-nm LED

- **LED element**
 - 0.1-mm x 0.1-mm
 - CW 0.15-mW @ 20 mA
 - Pulsed 0.45-mW @ 70 mA
- **Small LED array**
 - 2x2 in 0.3-mm x 0.3 mm
 - CW 0.6-mW @ 100 mA
 - Pulsed 1.8-mW @ 150 mA
- **Large LED array**
 - 5x5 in 1-mm x 1 mm
 - CW 3.75-mW @ 100 mA
 - Pulsed 11.25-mW @ 150 mA

340-nm LED

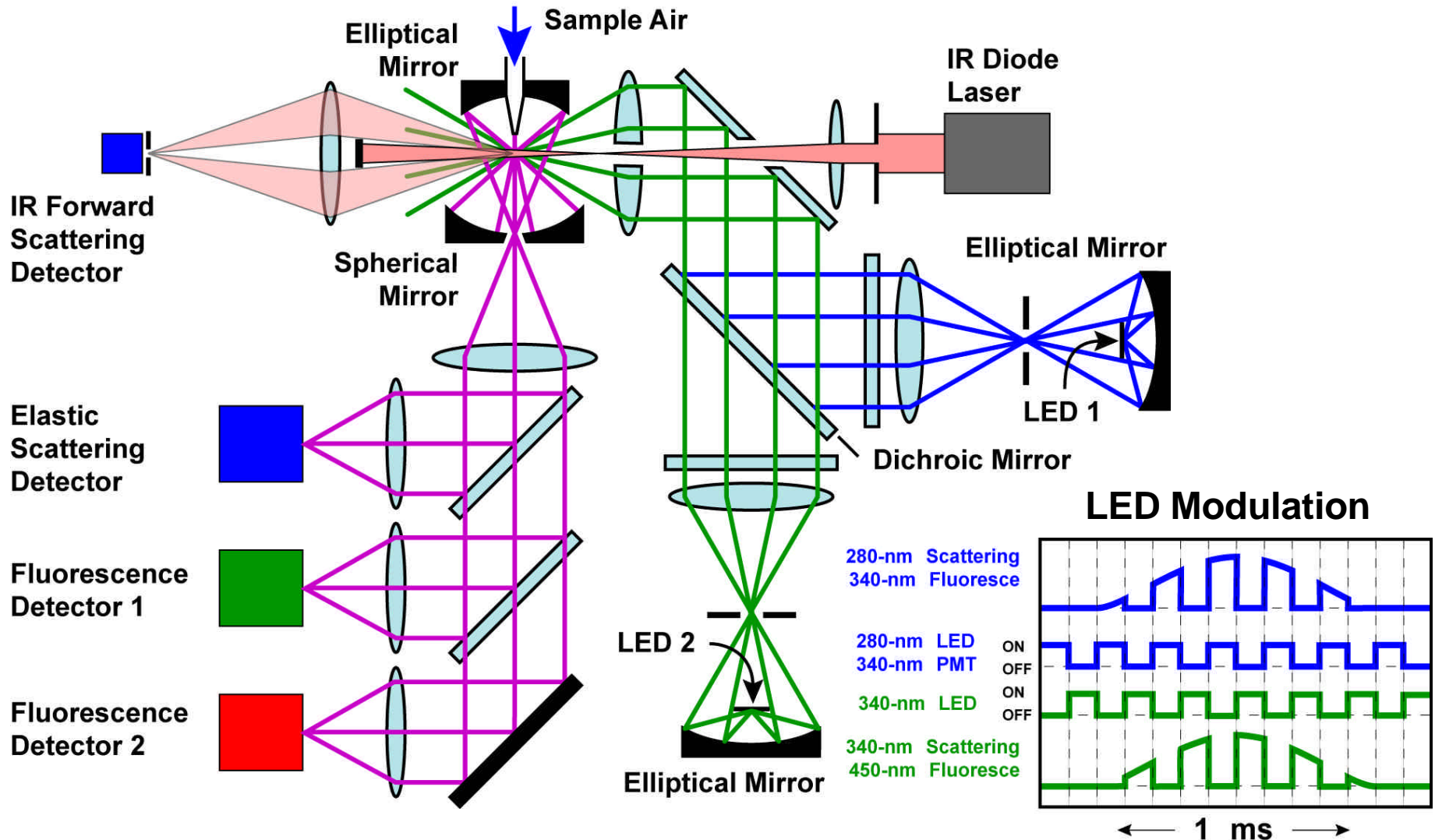
- **Small LEDs**
 - 0.3-mm x 0.3-mm
 - CW 0.8-mW @ 20 mA
 - Pulsed 8-mW @ 200 mA
- **Large LEDs**
 - 0.9-mm x 0.9 mm
 - CW 2.75-mW @ 100 mA
 - Pulsed 23-mW @ 1000 mA

All LEDs mounted on TO-46 headers.

Pulsed operation is for 1-ms pulse duration at 100-Hz repetition rate.

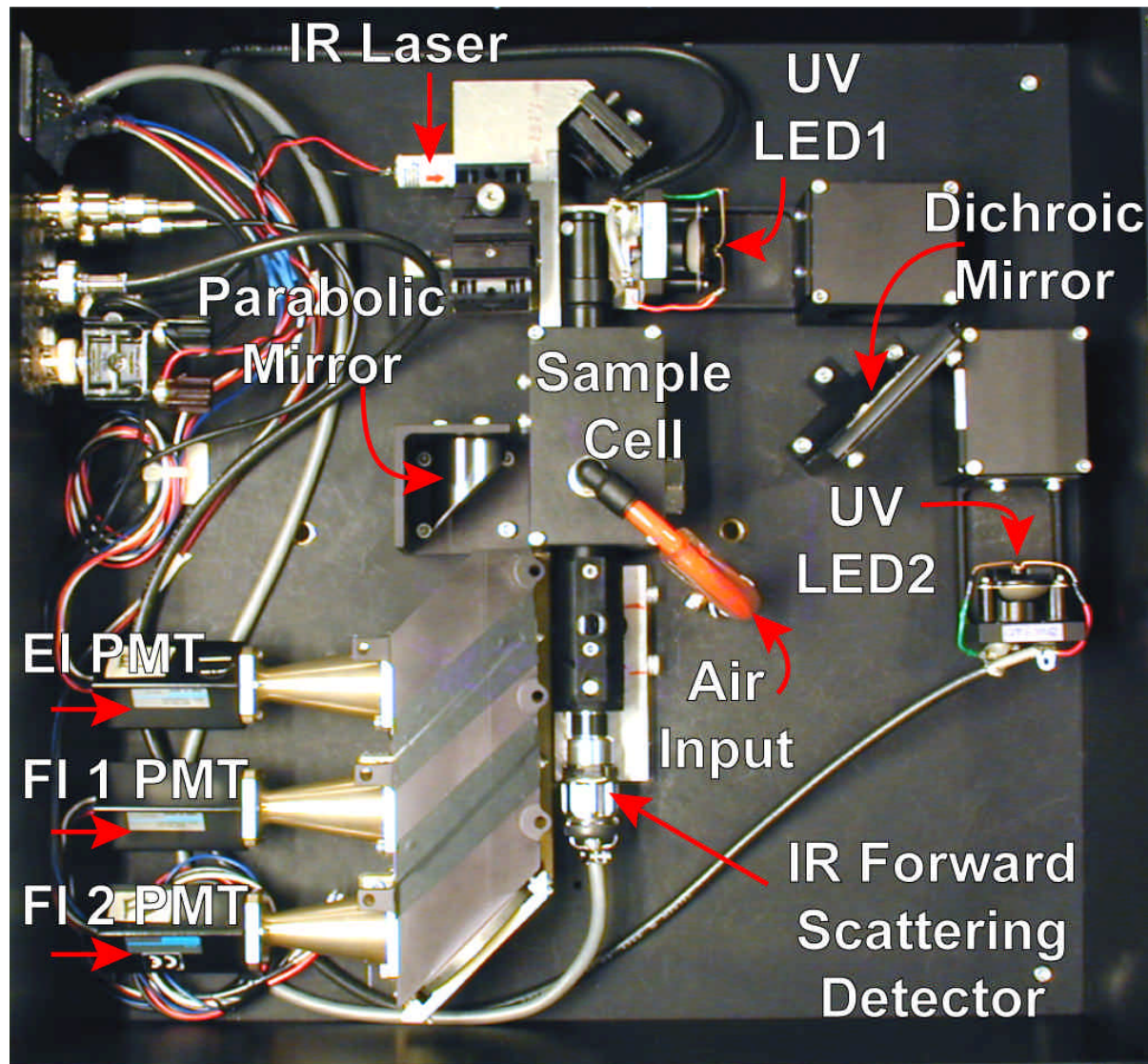


Optics Concept



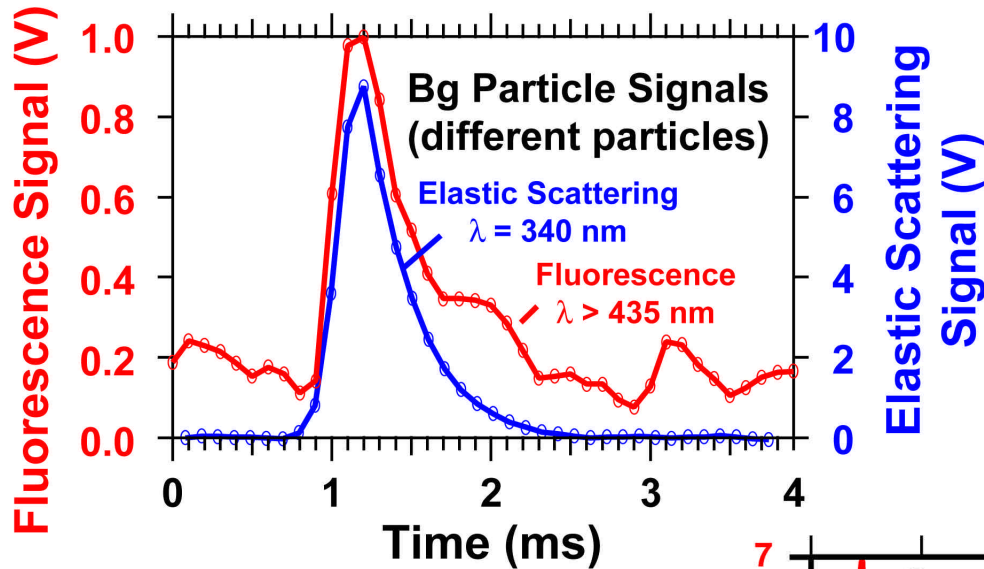


Proof-of-Concept Apparatus





Proof-of-Concept Fluorescence Data

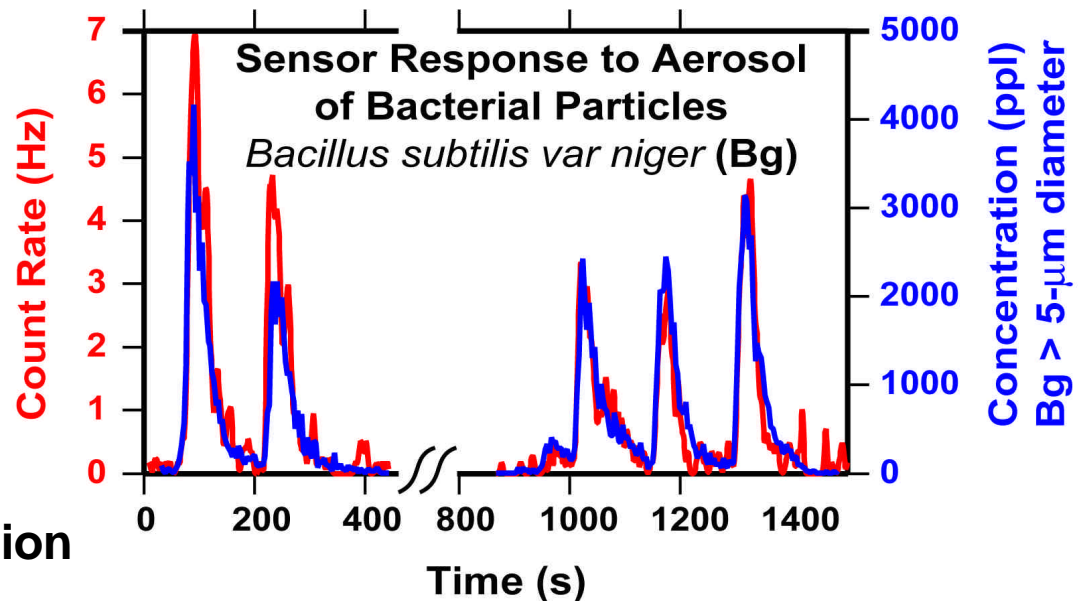


Detection of aerosolized single particle biological agent simulants (Bg) using 340-nm Cree LED

Responsivity

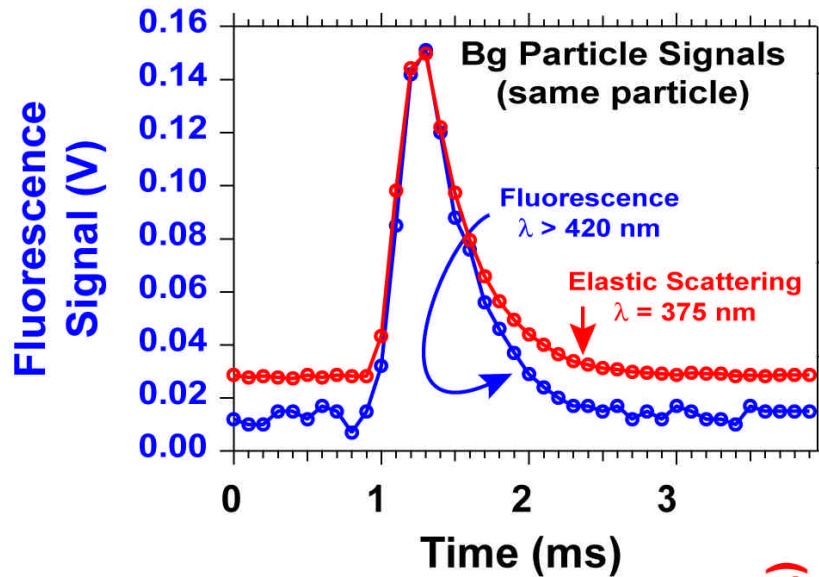
1 Hz count rate

700 ppl concentration





Proof-of-Concept Fluorescence Data



Elastic Scattering Signal (V)

Detection of aerosolized single particle biological agent simulants (Bg) using 375-nm LED

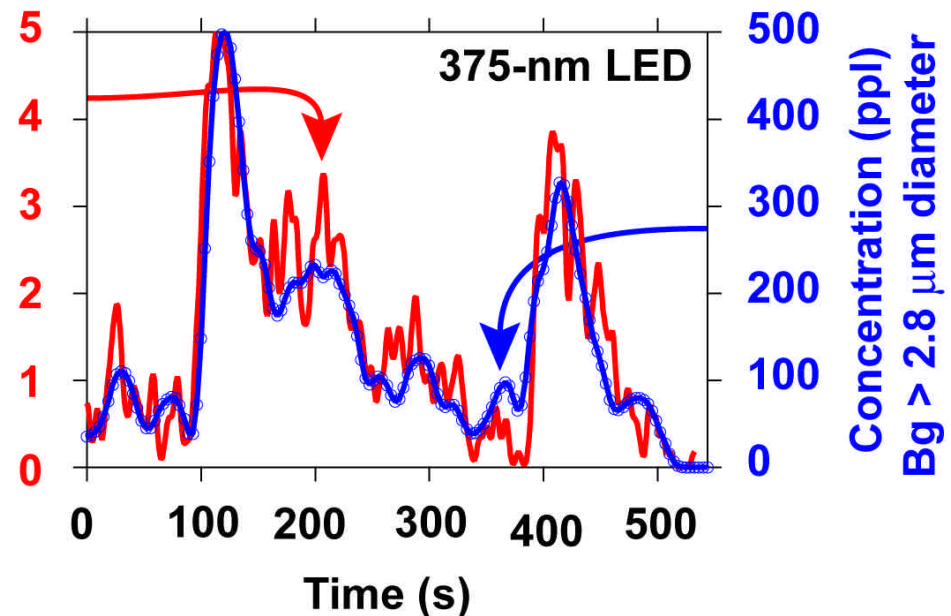
Responsivity

1 Hz count rate



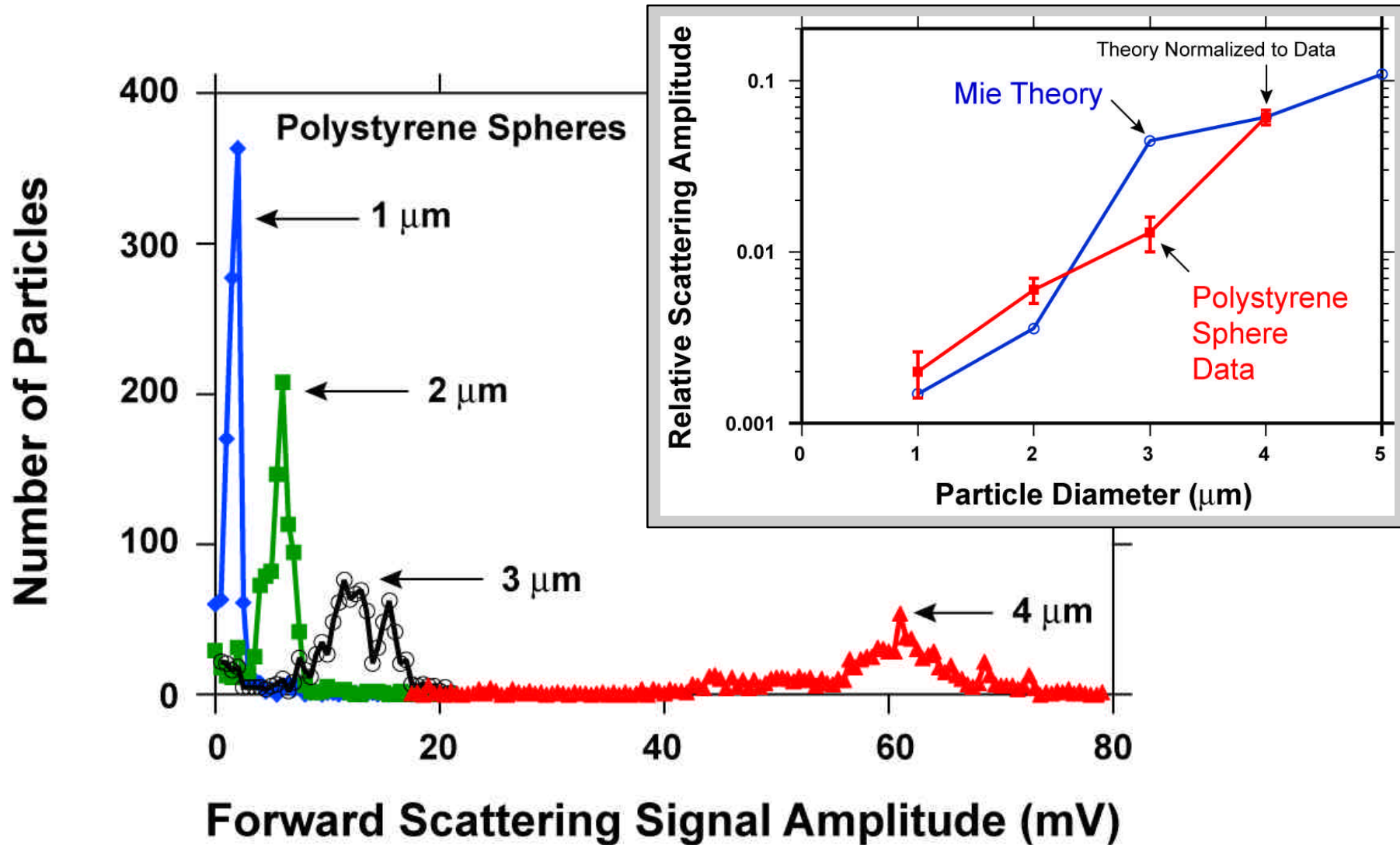
100 ppl concentration

Count Rate (Hz)





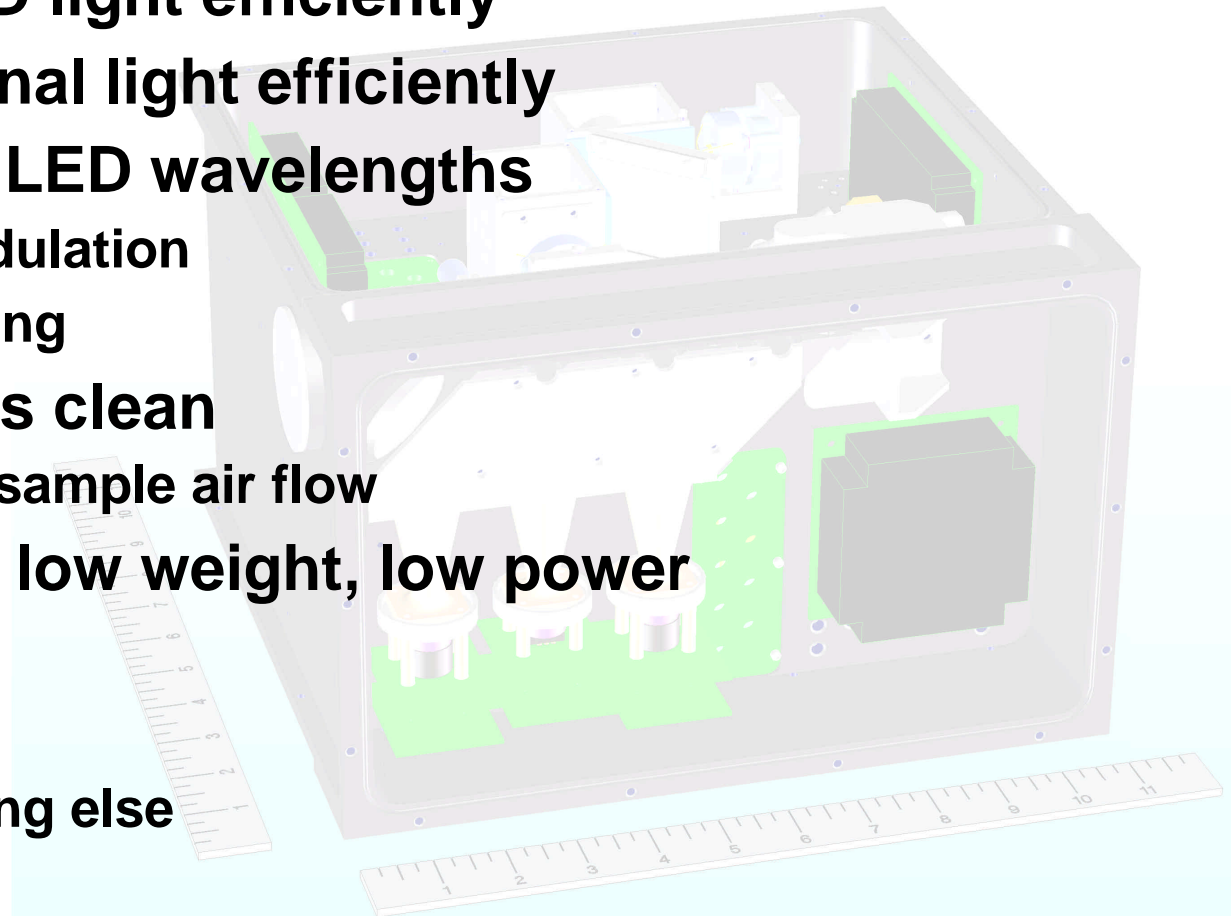
Proof-of-Concept Particle-Sizing Data





Engineering Development Unit: Challenges

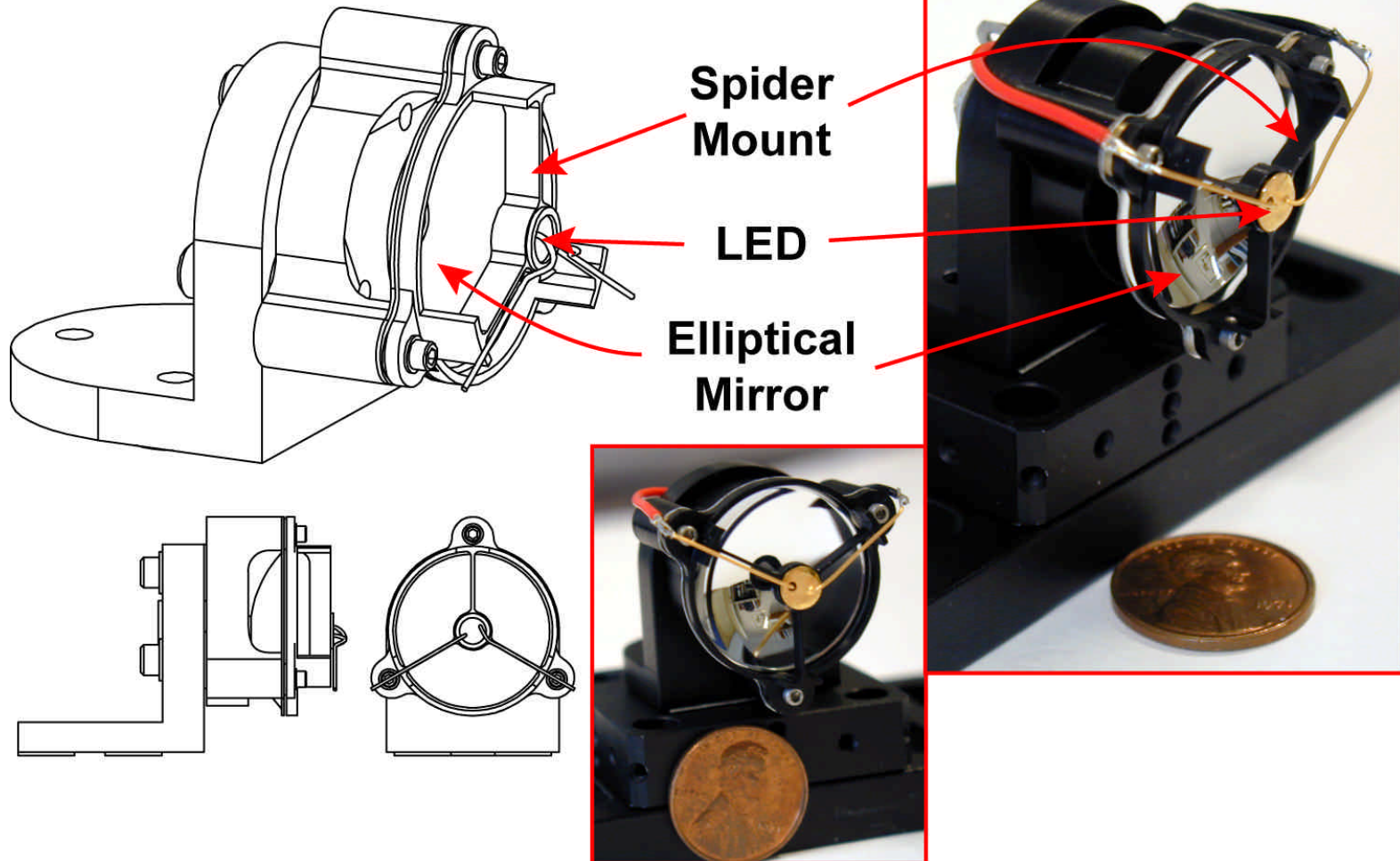
- **Collect LED light efficiently**
- **Collect signal light efficiently**
- **Utilize two LED wavelengths**
 - LED modulation
 - PMT gating
- **Keep optics clean**
 - Confine sample air flow
- **Small size, low weight, low power**
- **Low Cost**
 - LEDs
 - Everything else





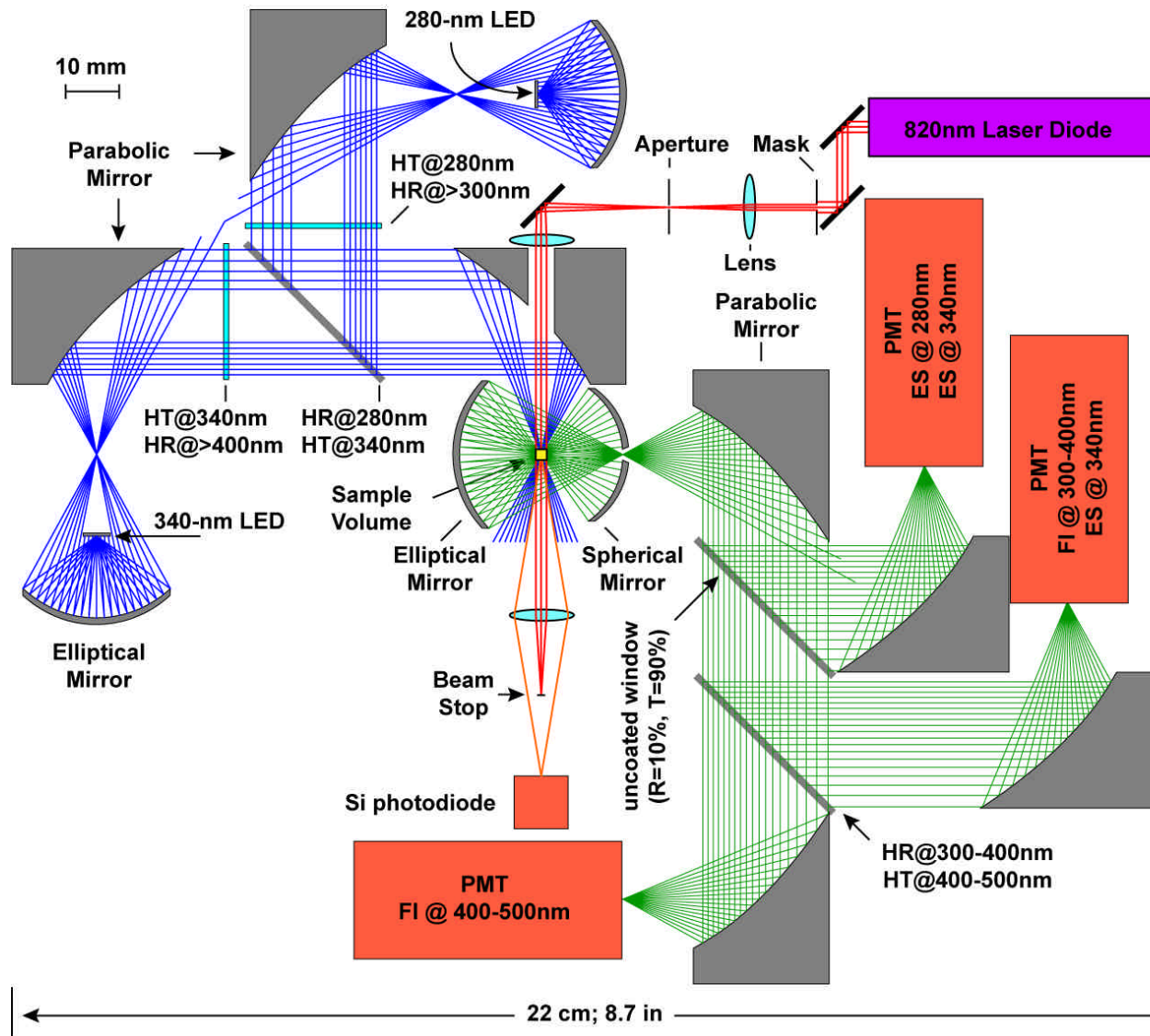
Engineering Development Unit Design: Optics

LED Mount and Elliptical Mirror





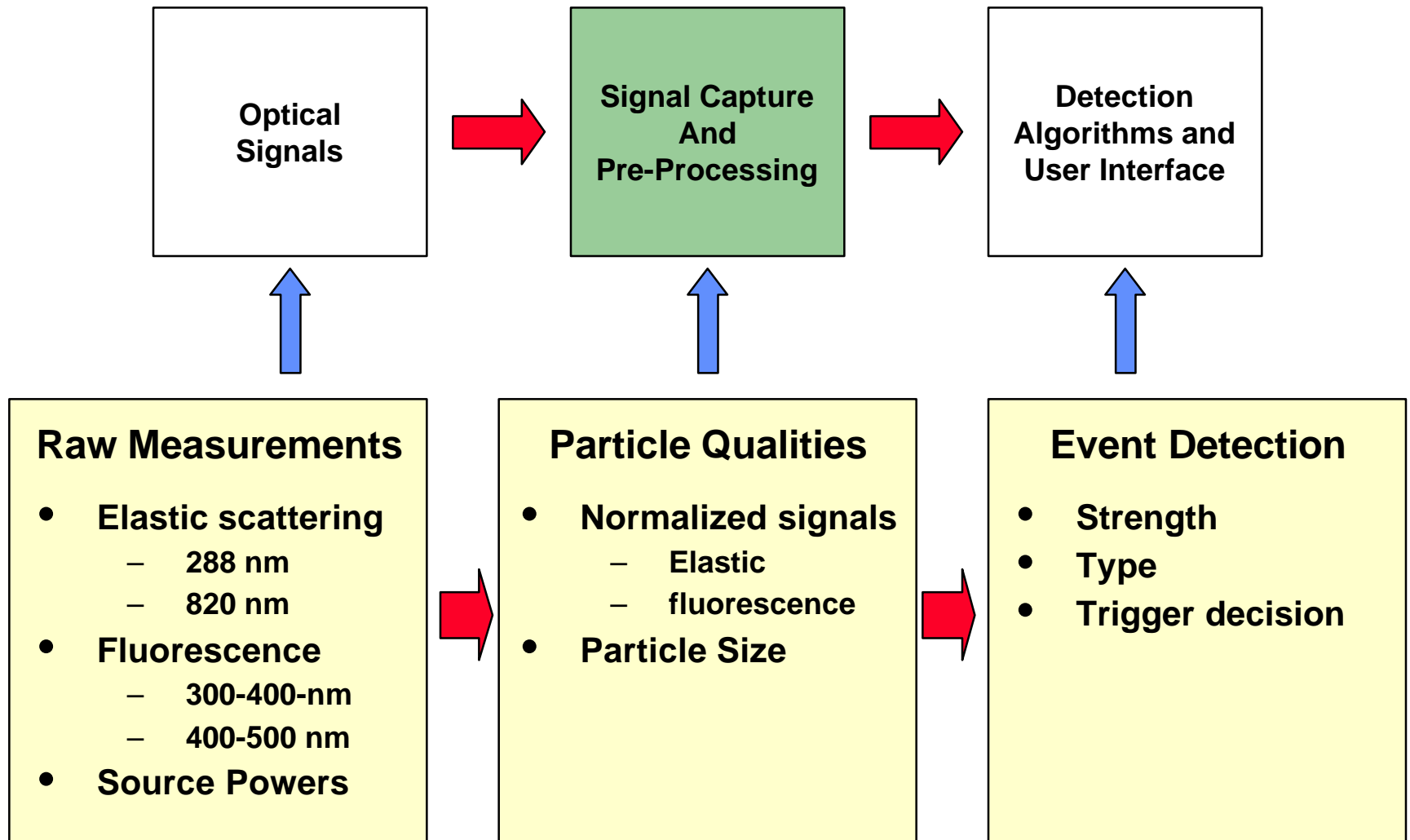
Engineering Development Unit Design: Optics





Engineering Development Unit Design: Electronics

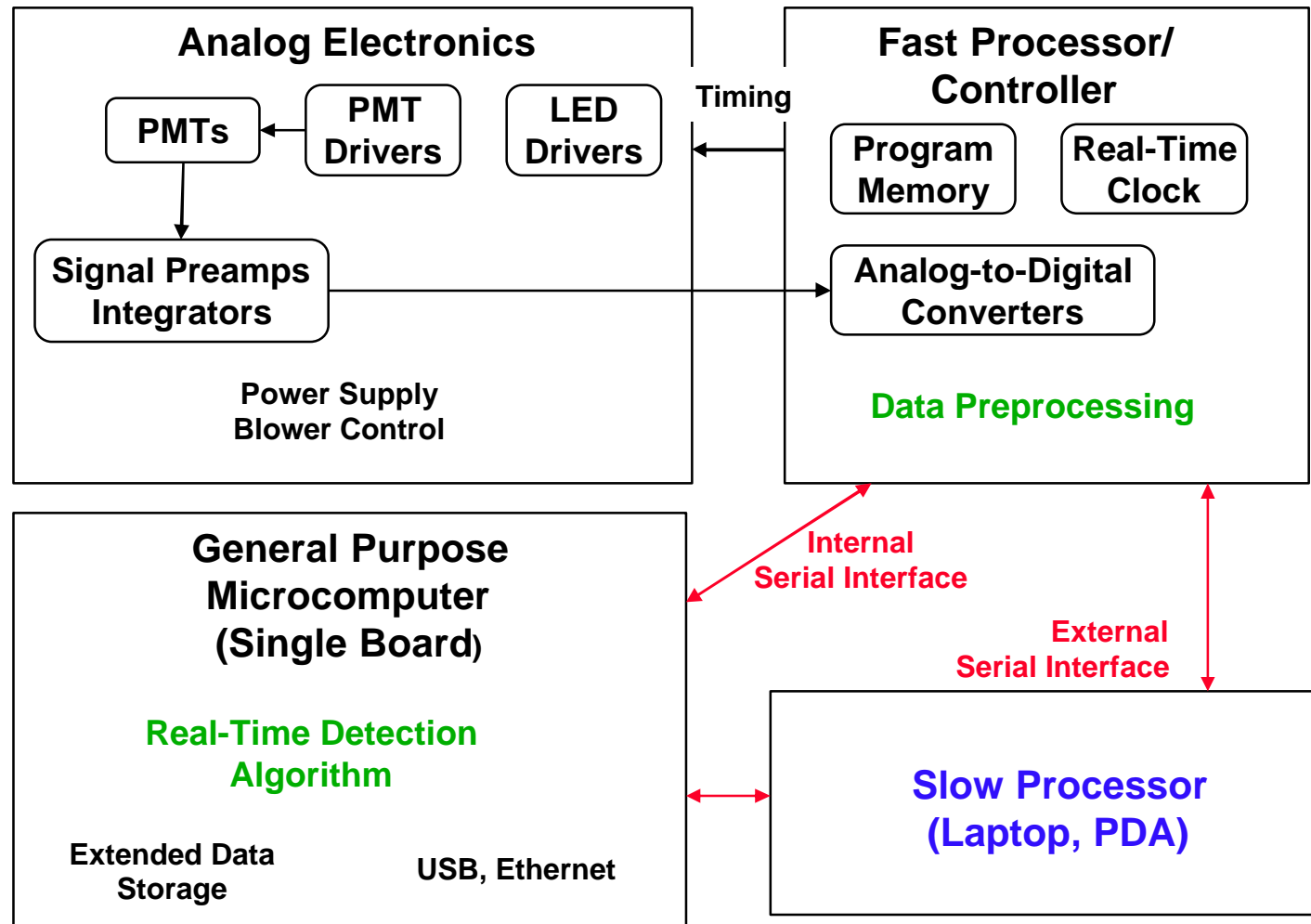
Electronics Functional Concept





Engineering Development Unit Design: Electronics

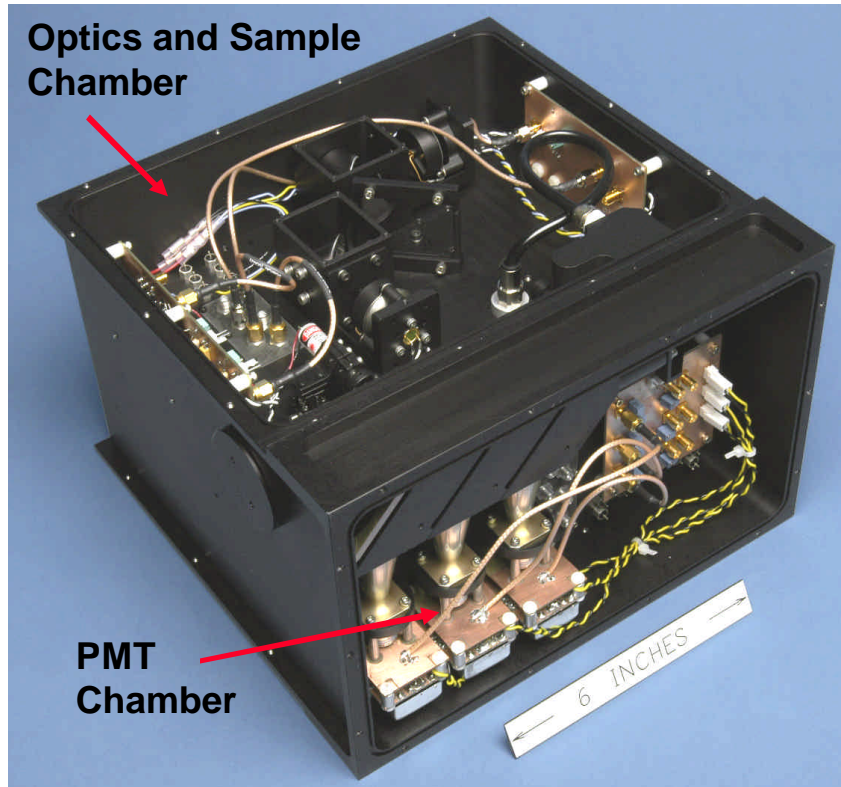
Electronics Block Diagram



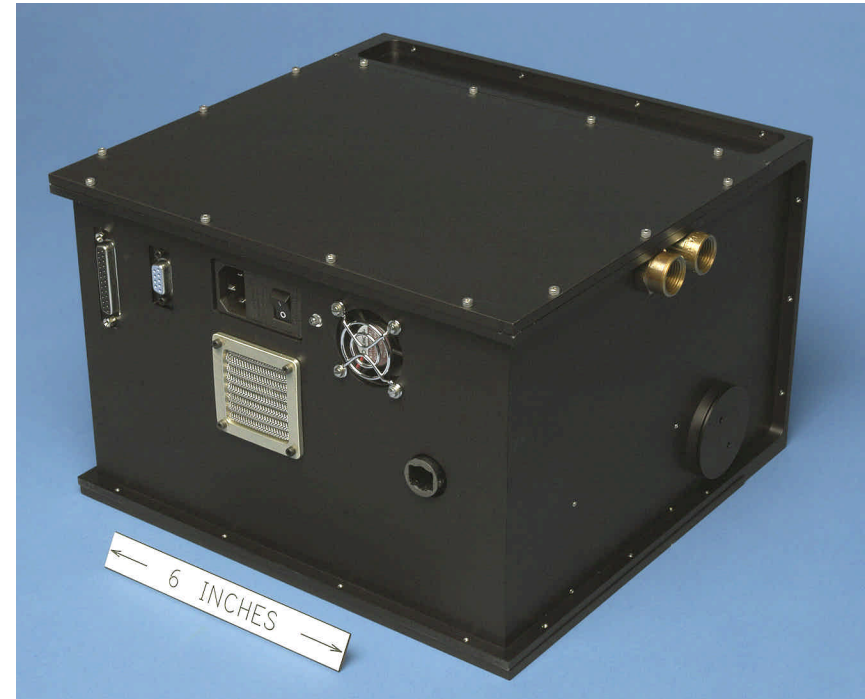


Engineering Development Unit Design: Mechanical

Covers Off



Covers On



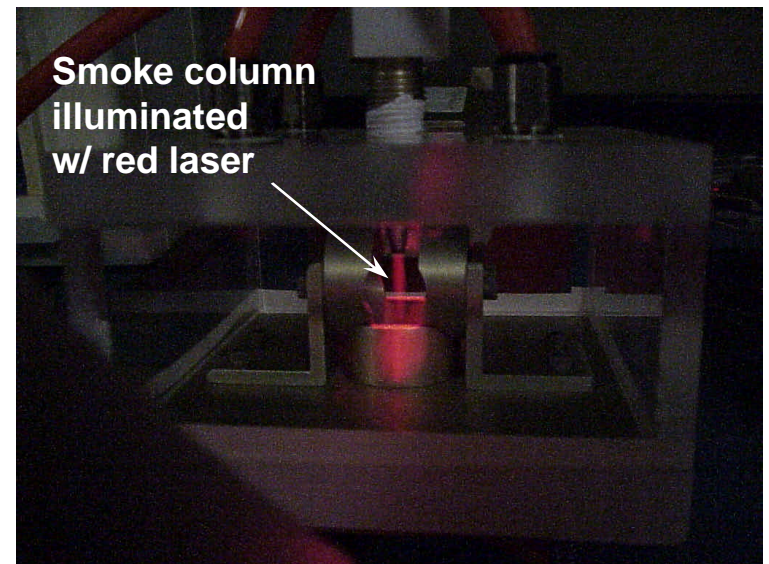
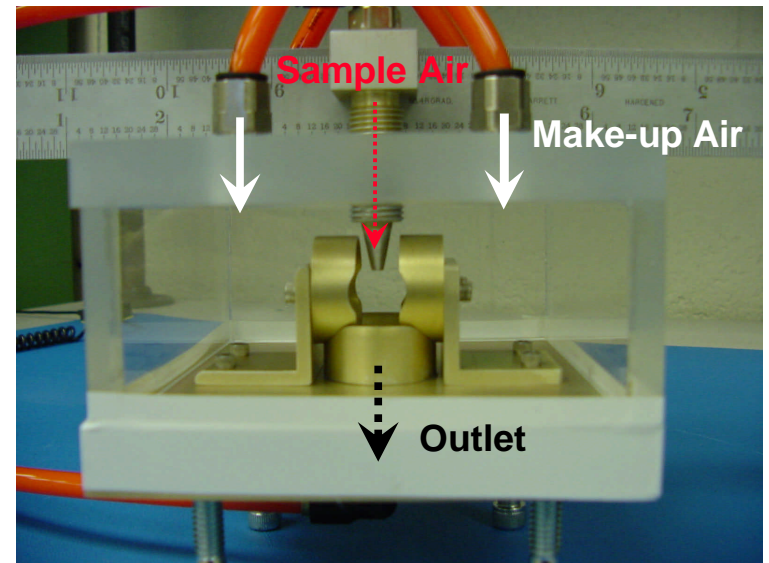
Compact design
Three compartments
Easy access to all compartments

Weight approx. 17.5 lbs
11.1" x 10.9" x 7.3"
Can be used in any orientation



Engineering Development Unit Design: Sample Air Handling

- Sample air contained in a free jet
- Additional make-up air confines sample air and prevents optics contamination
 - HEPA filtered make-up air
- Laminar flow
 - Straight particle trajectories
 - Well defined air speed
- Test apparatus constructed
 - Confirmed that make-up air isolates sample air





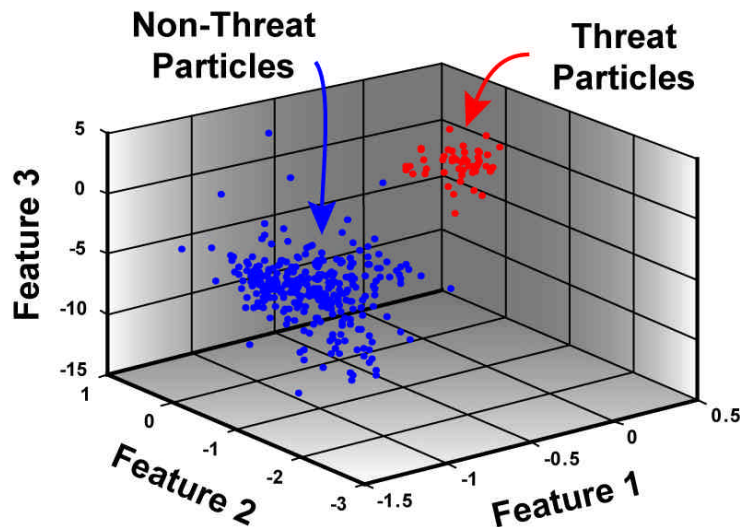
Engineering Development Unit Design: Algorithm

1. Training

- Learn background spectral signature in multidimensional space
 - Particle size
 - Fluorescence
 - Elastic scattering
 - Combinations of features

2. Change Detection

- Compare present aerosol spectral signature with background signature
 - Significant difference indicates new type of aerosol



3. Discrimination

- Determine if event occurred in threat region
 - Generate trigger event
 - Track multiple concurrent events
 - Report event strengths



Summary

- **Analysis of LED based biodetection**
- **Design and construction of three test-bed generations**
- **First demonstration of LED based detection of biological agent simulants (*Bacillus subtilis var niger (Bg)*, *ovalbumin*)**
- **Measurement of elastic and fluorescence aerosol signals using 280-nm and 340-nm LEDs**
 - Bg, Ovalbumin, Dirt
 - > 3-mm Bg particle size
 - > 1 Hz per 100 - 700 ppl Bg responsivity
- **Design and construction of engineering development unit**



Biological Agent Sensor Testbed Performance Analysis

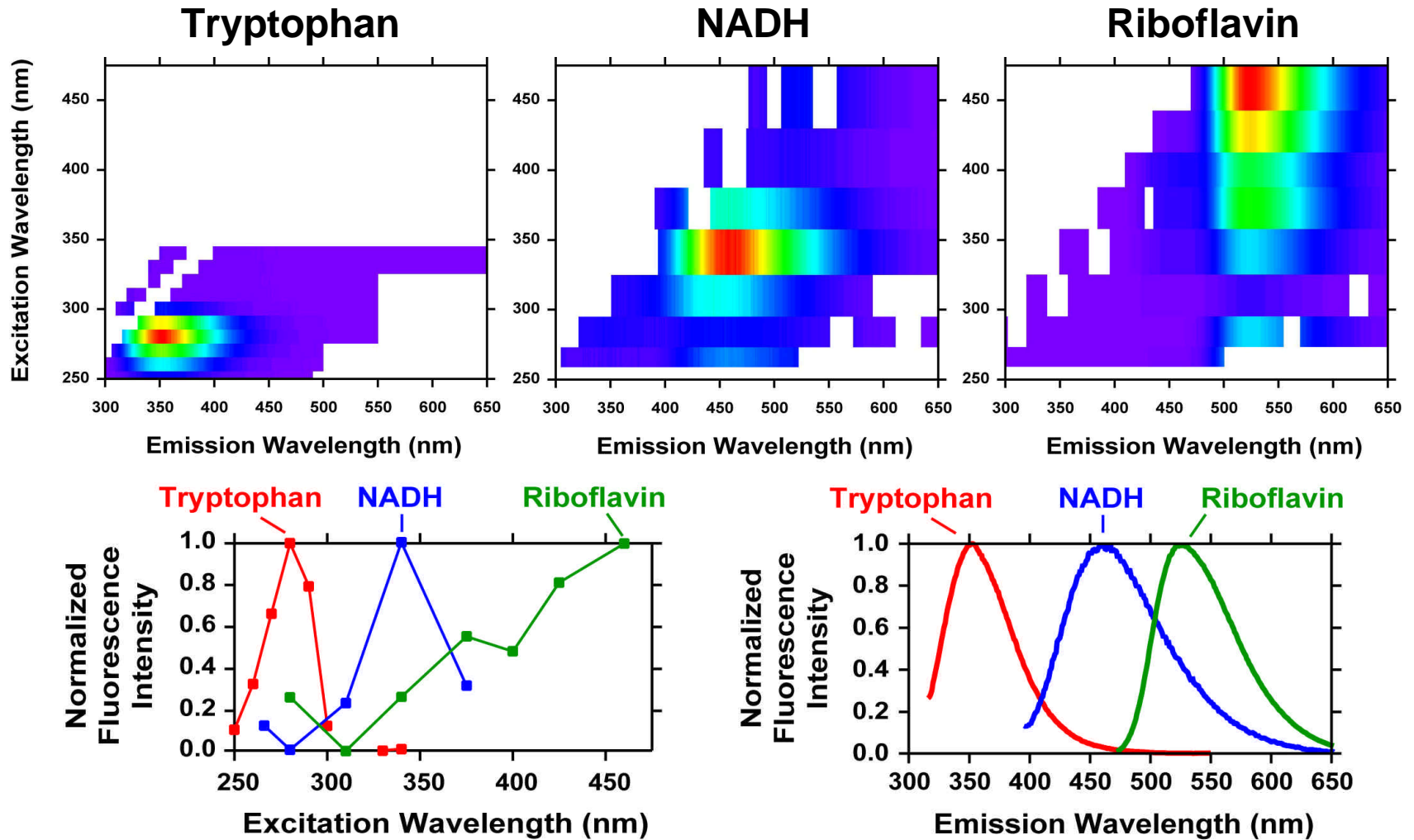


Performance Analysis

- **What LED wavelengths are optimal?**
 - Detection
 - Discrimination
- **What is the minimum detectable particle concentration?**
 - LED power dependence
 - LED wavelength dependence
- **Can particle size be determined from forward scattering?**
 - Wavelength
 - Angular range of integration
- **What happens at very high concentrations?**



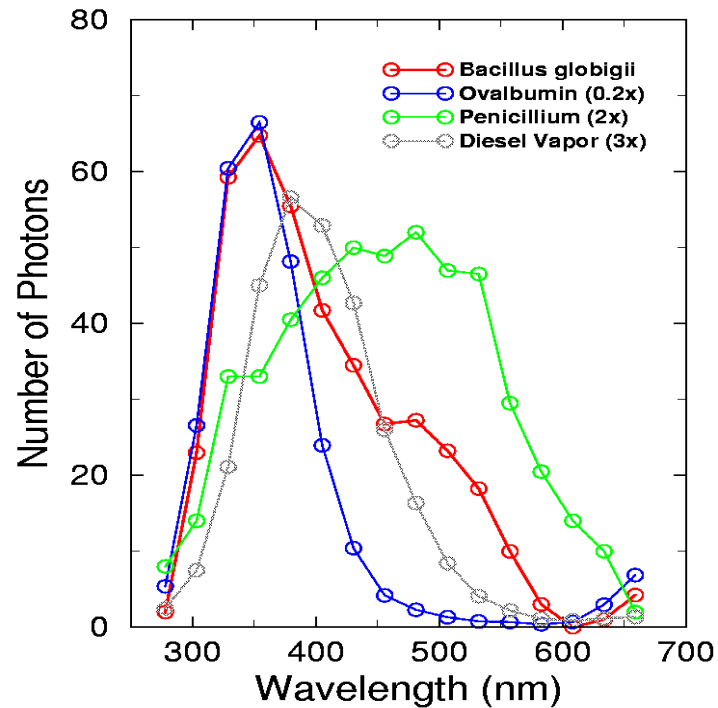
Bio-Fluorescence Spectra



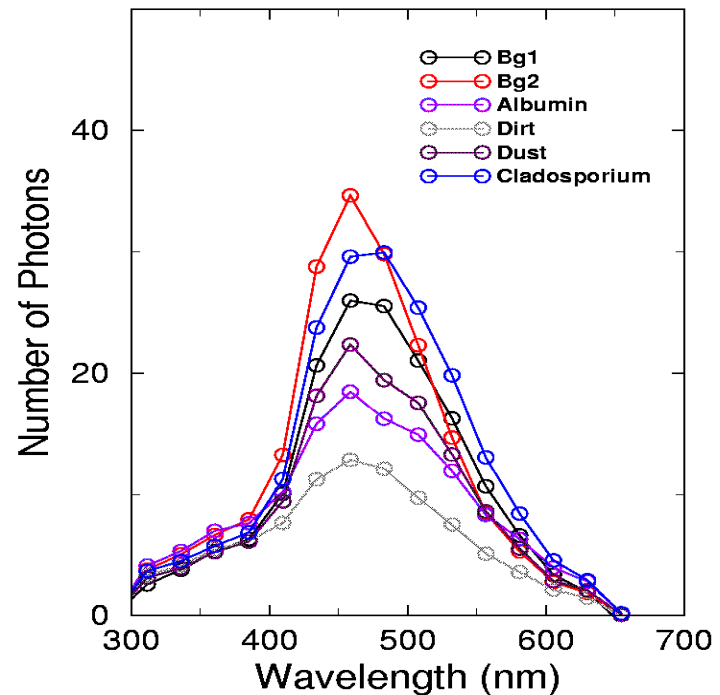


Fluorescence Spectra

266-nm Excitation



355-nm Excitation



266-nm spectra provides more discrimination than 355 nm spectra.



Performance Equations

of detected photons
↓

light power
↓

illumination time
↙

light wavelength
↘

$$N_p = h_c h_d \frac{P t l}{A h c} S$$

collection efficiency
↗

detection efficiency
↗

light cross-sectional area
↑

Planck's constant
↖

speed of light
↖

cross-section
←

of detected particles
↓

$$n = f C T$$

sample rate
↑

particle concentration
↗

detection time
←

sample cross-sectional area
↓

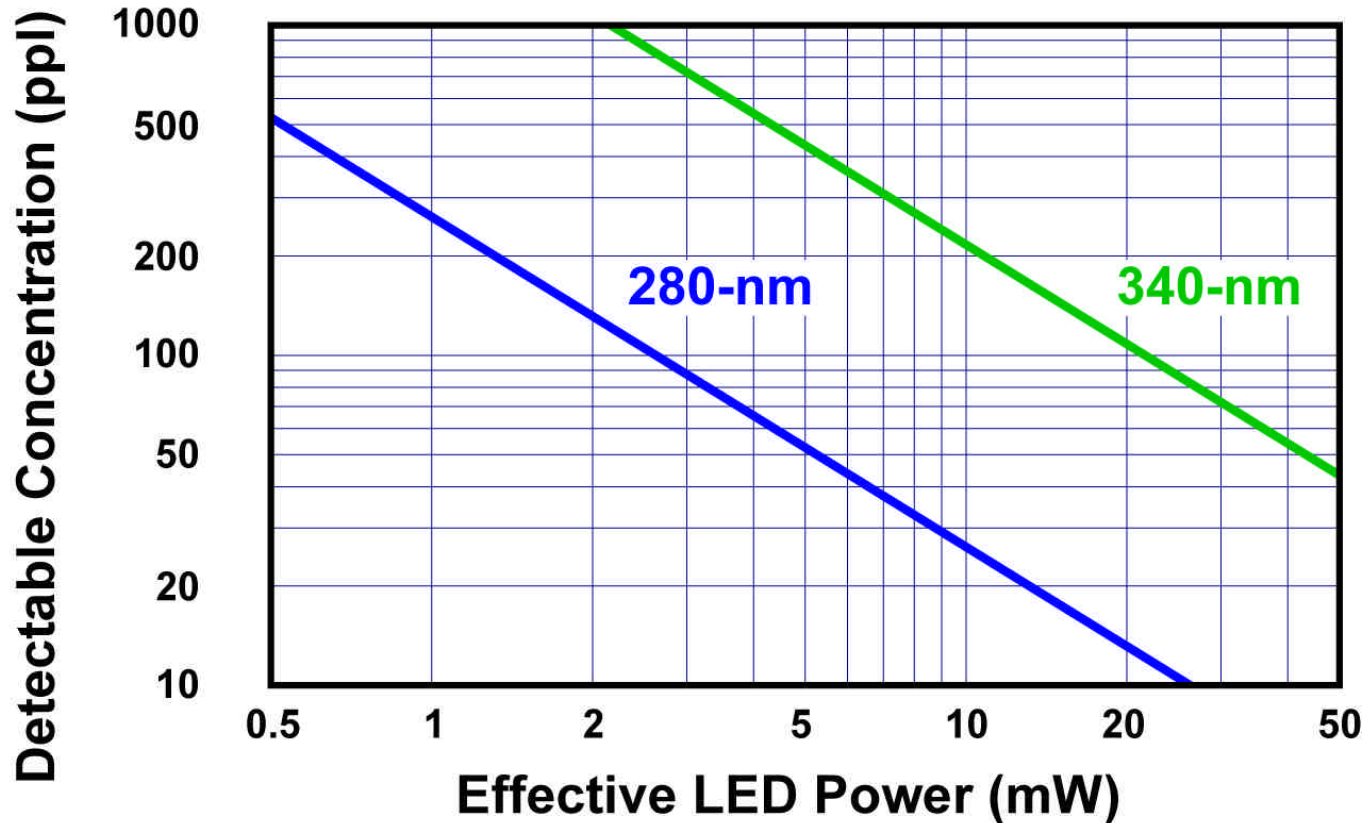
$$f = \frac{A L}{t}$$

sample length
←

sample transit time
←



Fluorescence Detection Analysis



| | |
|---|--|
| Number of detected particles (N_b) | 100 particles |
| Particle fluorescence cross-section (σ) | $(50, 5) \times 10^{-12} \text{ cm}^2$ @ (280, 340) nm |
| Number of detected photo-electrons (N_{pe}) | 100 photoelectrons |
| Photon collection, detection efficiency (h_d) | 30% , 15% |
| Threat detection time (T) | 60 s |



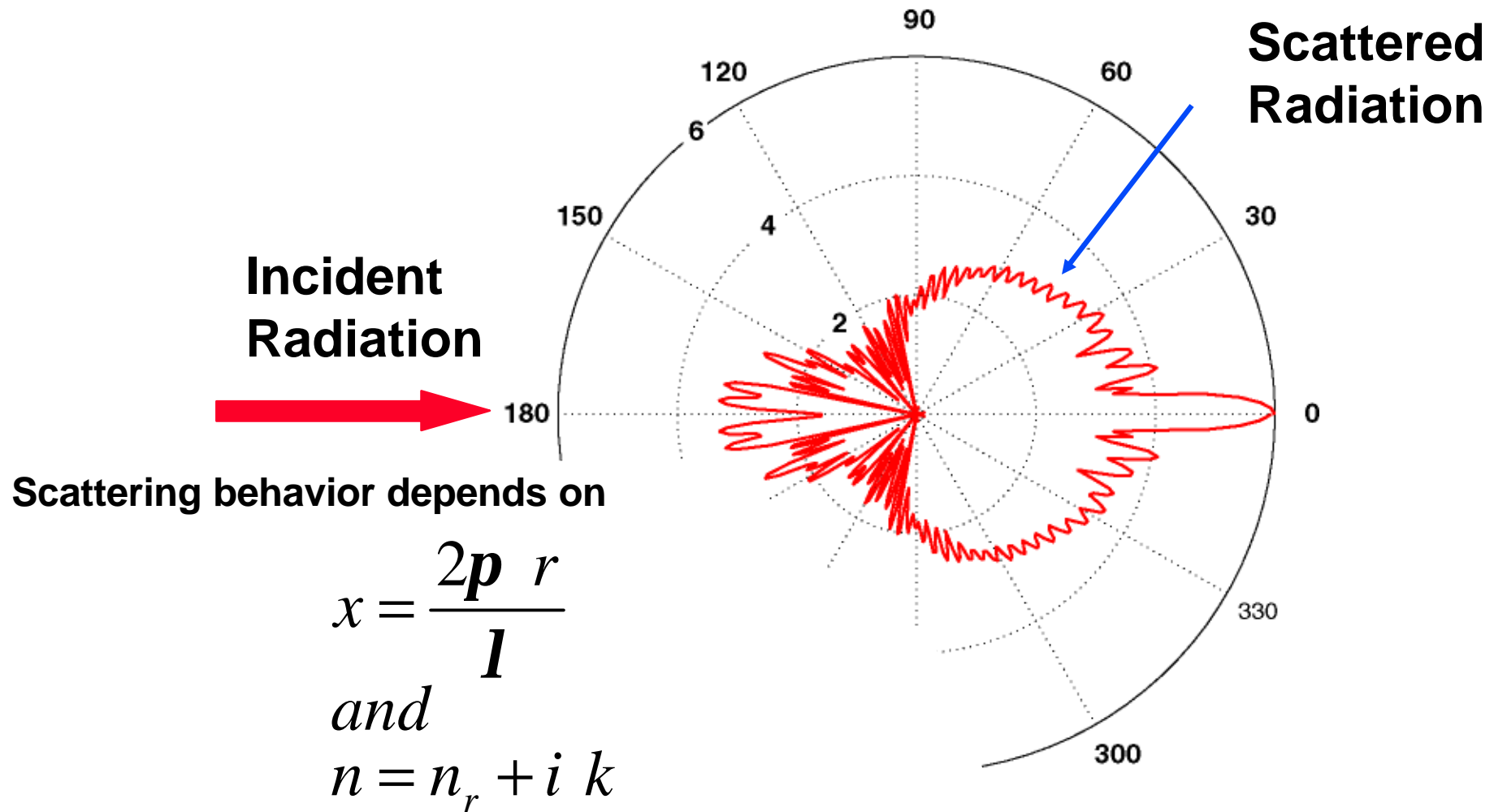
Example Calculation; 1-mm diameter LEDs

| | elastic scattering | fluorescence | fluorescence |
|--|--------------------|--------------|--------------|
| LED wavelength (cm) | 2.80E-05 | 2.80E-05 | 3.40E-05 |
| optical cross section (cm ²) | 5.00E-09 | 5.00E-11 | 5.00E-12 |
| LED diameter (cm) | 1.00E-01 | | |
| sample cross-sectional area (cm ²) | 4.00E-02 | | |
| Total Pulsed LED Power (W) | 3.75E-03 | 3.75E-03 | 2.30E-02 |
| Effective Pulsed LED power (W) | 1.44E-03 | 1.44E-03 | 8.81E-03 |
| photon collection efficiency | 3.00E-01 | | |
| photon detection efficiency | 1.50E-01 | | |
| illumination time, τ (s) | 1.20E-03 | | |
| detected photons | 1.36E+04 | 1.36E+02 | 1.02E+02 |
| concentration uncertainty ($\Delta C/C$) | 1.00E-01 | | |
| number of detected particles | 1.00E+02 | | |
| detection time (min) | 1.00E+00 | | |
| sample length (cm) | 2.00E-01 | | |
| sample rate (liter/min) | 4.00E-01 | | |
| detectable concentration (ppl) | 2.50E+02 | | |

* **Sample volume = (2 x LED diameter)³ = 8 x 10⁻⁶ liters**



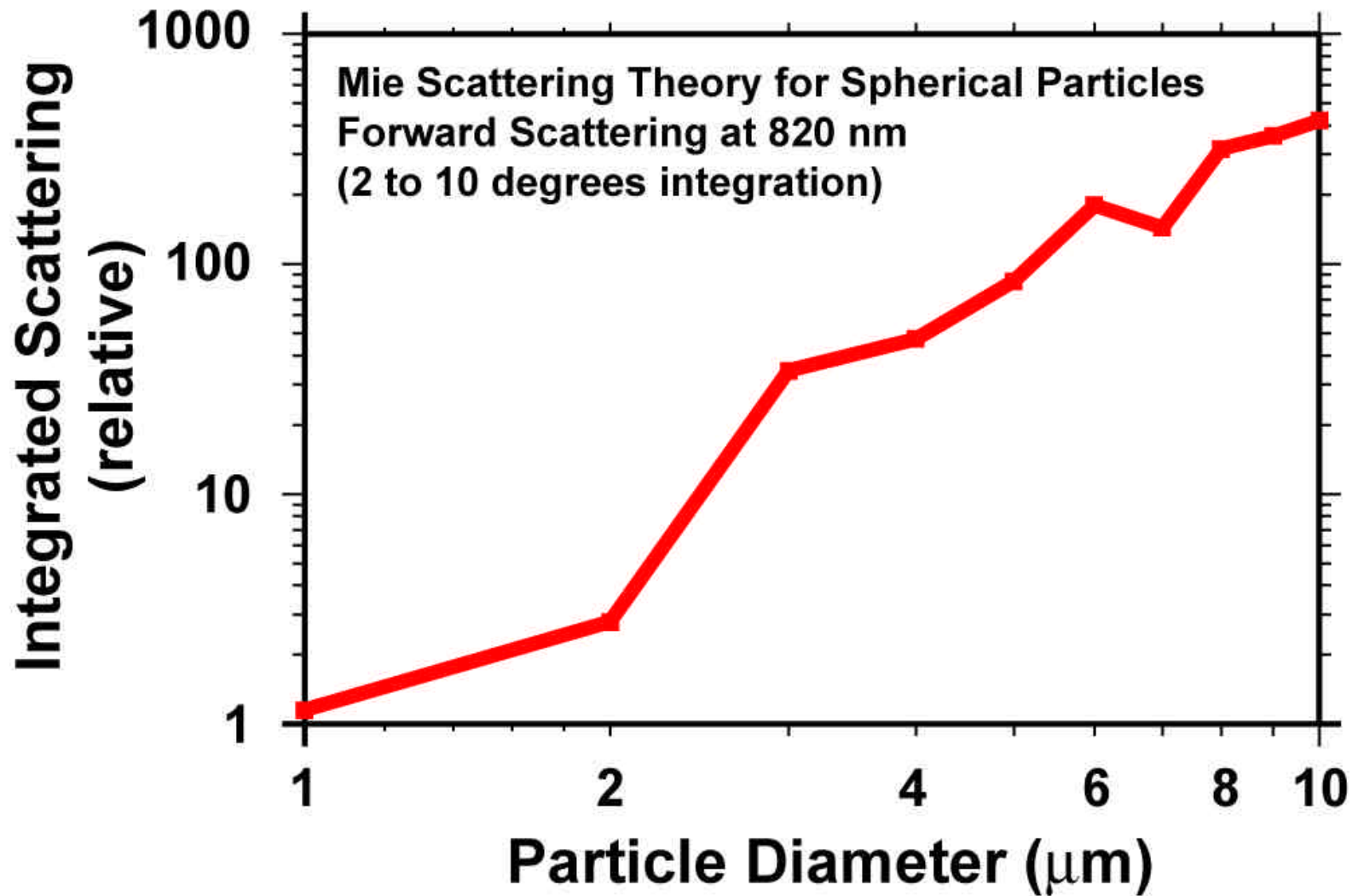
Elastic Scattering Angular Distribution



Barnett, D., The Mie scattering software is available at
<http://www.lboro.ac.uk/departments/el/research/optics/matmie/mfiles.html> (May 2002).



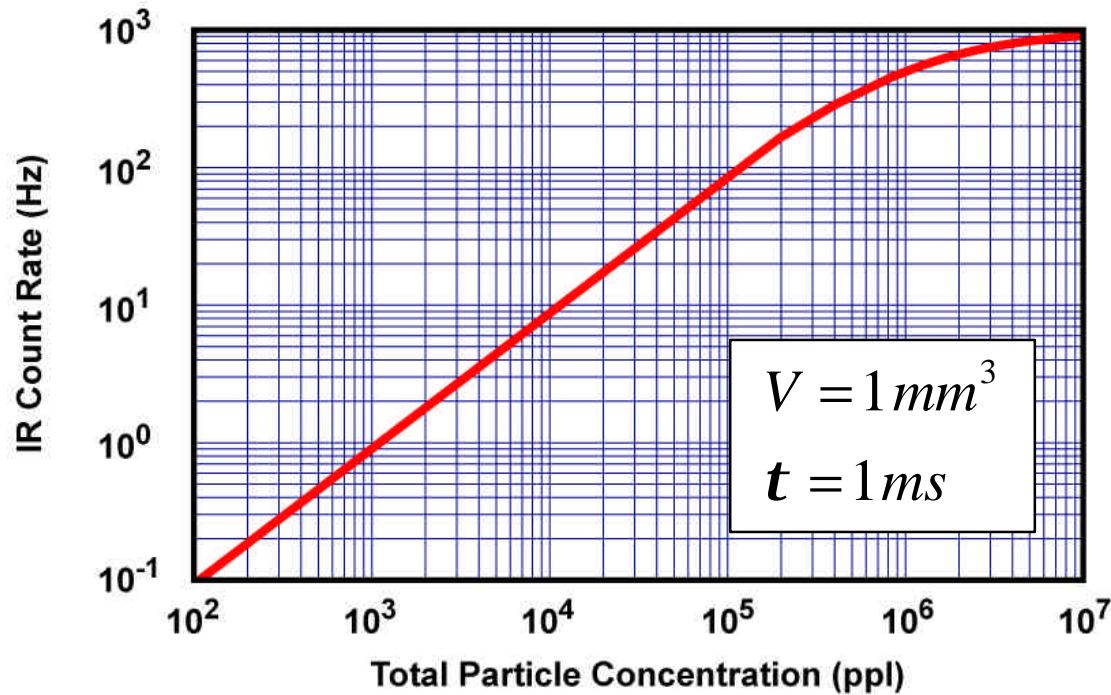
Elastic Scattering and Particle Size





IR Particle Count Rate

$$f_{IR} = \frac{\Phi C}{1 + \frac{\Phi C}{f_{IR_max}}} = \frac{1}{t} \left(\frac{V C}{1 + V C} \right); \text{ where } \Phi = \frac{V}{t} \text{ and } f_{IR_max} = \frac{1}{t}$$

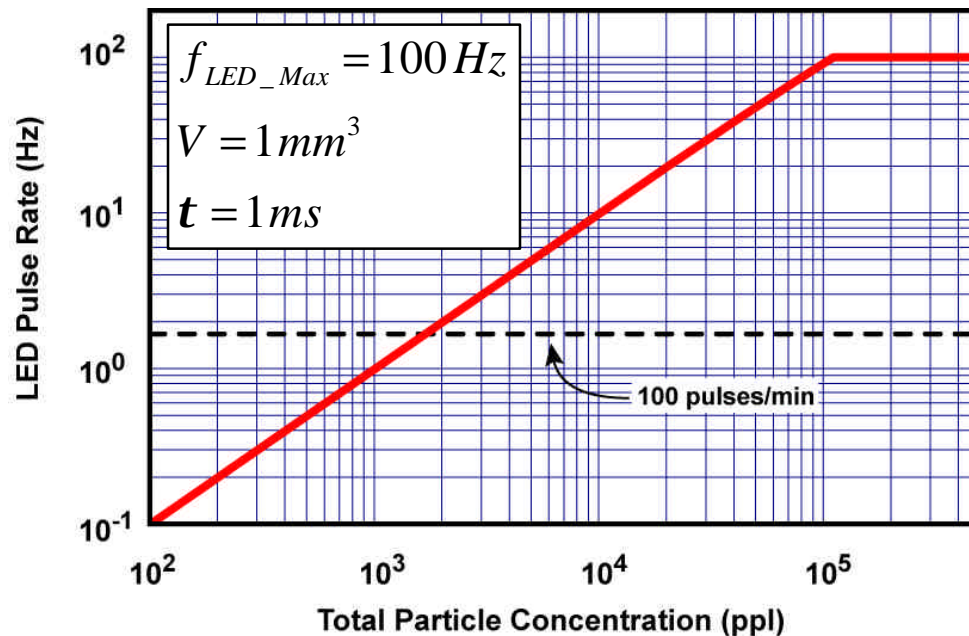




LED Pulse Rate

$$f_{LED} = f_{IR} \quad ; f_{IR} < f_{LED_Max}$$

$$f_{LED} = f_{IR_Max} \quad ; f_{IR} \geq f_{LED_Max}$$





Threat Detection

$$f_{threat} = f_{LED} \frac{C_{threat}}{C}$$

C_{threat} = threat concentration

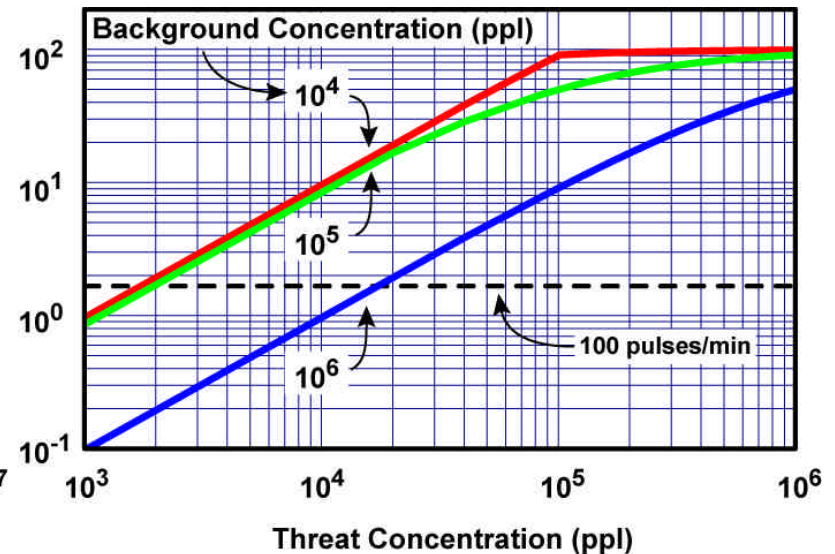
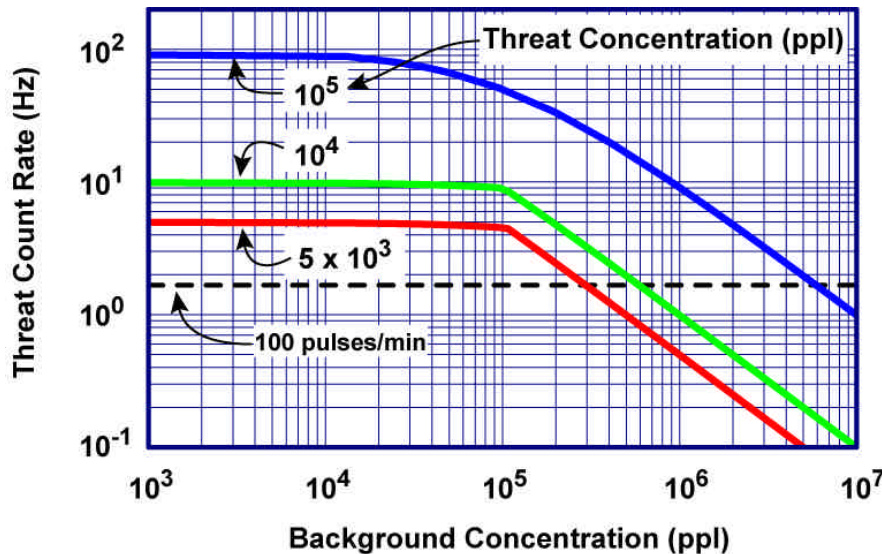
$C_{nonthreat}$ = nonthreat concentration

$C = C_{threat} + C_{nonthreat}$ = total concentration.

$$f_{LED_Max} = 100Hz$$

$$V = 1mm^3$$

$$t = 1ms$$





Minimum Detectable Threat Concentration

$$C_{threat_Min} = C_1 \text{ for } f_{LED} < f_{LED_Max}$$

$$C_{threat_Min} = C_2 \text{ for } f_{LED} = f_{LED_Max}$$

or

$$C_{threat_Min} = \text{Max}[C_1, C_2]$$

where

$$C_1 = \frac{\frac{1}{V} + C_{nonthreat}}{\frac{1}{t f_{alarm}} - 1}$$

and

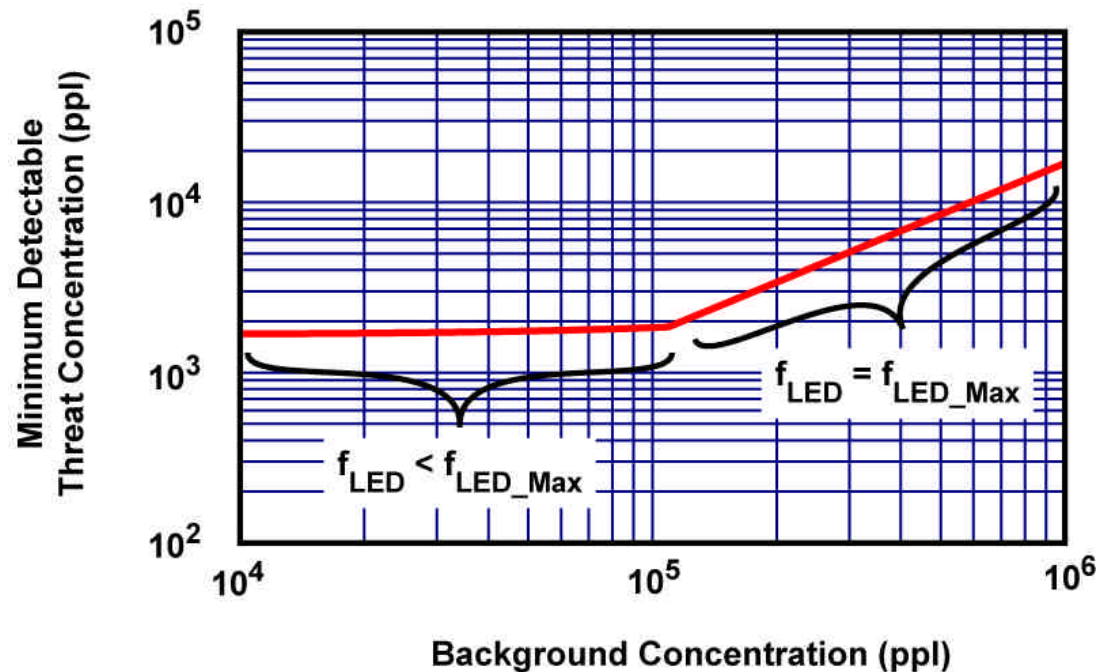
$$C_2 = \frac{C_{nonthreat}}{\frac{f_{LED_Max}}{f_{alarm}} - 1}$$

$$f_{alarm} = 1.667 \text{ Hz} = 100 / \text{min}$$

$$f_{LED_Max} = 100 \text{ Hz}$$

$$V = 1 \text{ mm}^3$$

$$t = 1 \text{ ms}$$

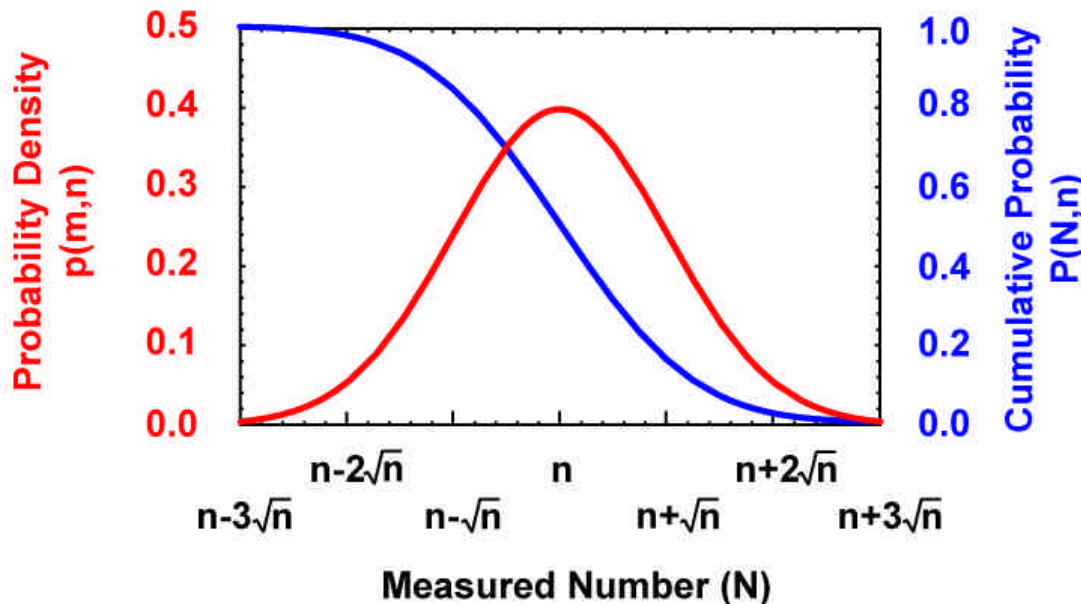




Detection Probability, P_d

Cumulative probability of counting N or more particles when the average count is n particles

$$P_d(N, n) = \int_N^{\infty} p(m, n) dm = \int_N^{\infty} \frac{e^{-\frac{(m-n)^2}{2n}}}{\sqrt{2\pi n}} dm \quad n \gg 1, m \gg 1$$

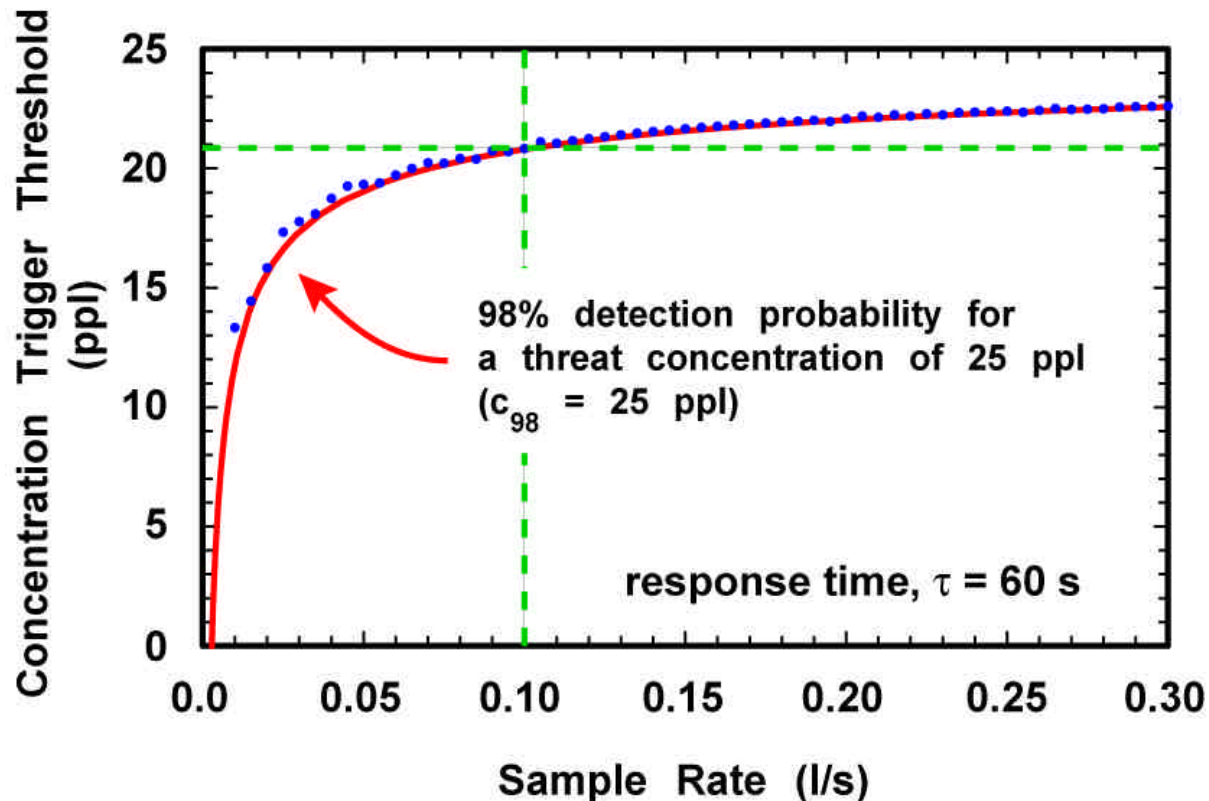


$$P_d(N, n) = 0.98, \text{ for } N = n - 2.05375\sqrt{n} \text{ where : } n \gg 1$$



Concentration Trigger Threshold

$$c_T = c_{98} \left(1 - \frac{2.05}{\sqrt{c_{98} f t}} \right)$$





Summary

- **This analysis shows that given currently available UV LEDs it should be possible to detect biological agent aerosols at concentrations of interest.**
- **This analysis also shows how particle triggering can result in nonlinear threat detection**
 - **It is also possible to use this analysis to correct the measured threat concentration using the background concentration**
- **High probability of detection requires setting the concentration detection threshold below the mean concentration of interest.**



Biological Agent Sensor Testbed Optical Design

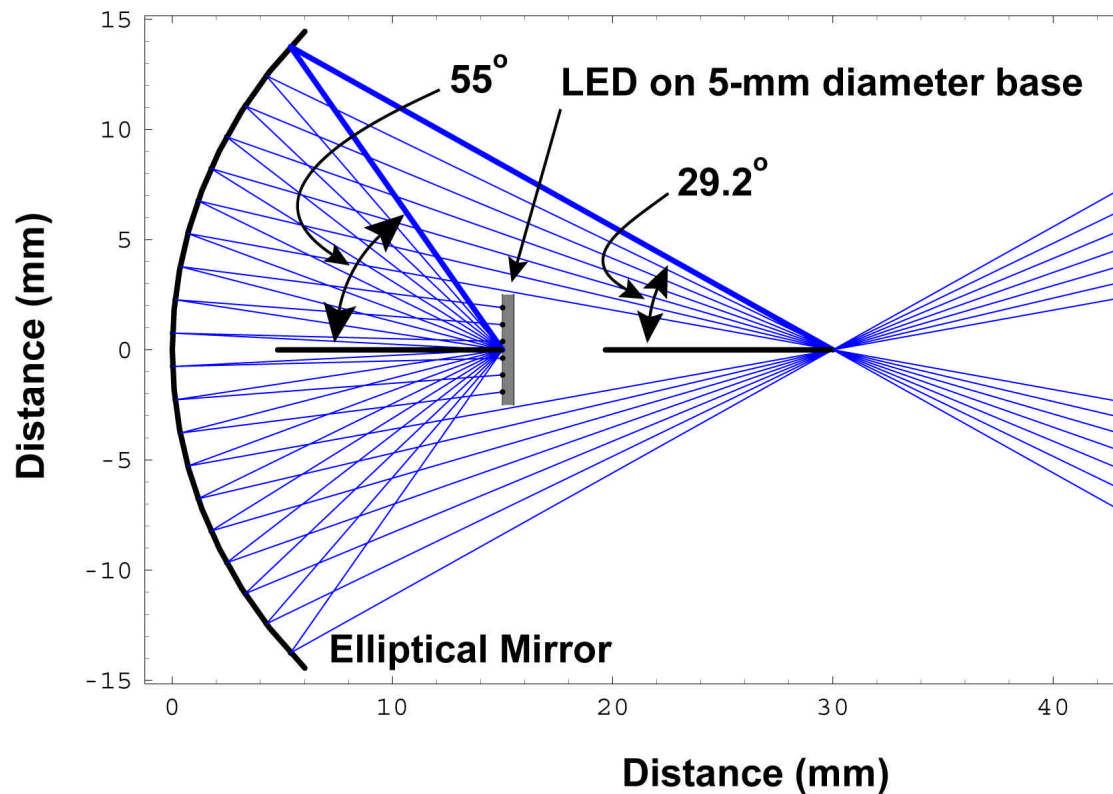


Challenges

- **Collect LED light efficiently**
 - Re-image LED light into sample volume
- **Mount LED with good heat sinking**
- **Collect signal light efficiently**
- **Utilize two LED wavelengths**
- **Measure particle size**
- **Avoid particle concentrator**
 - Need large sample volume
- **Keep optics clean**
- **Utilize low cost optical components and mounts**
 - Design optical system with minimum of optical adjustments



LED Light Collection Mirror

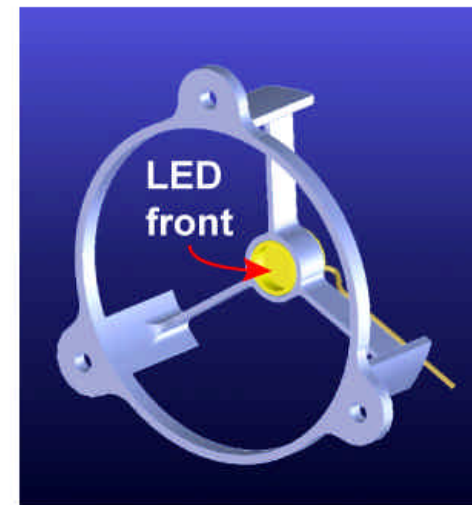
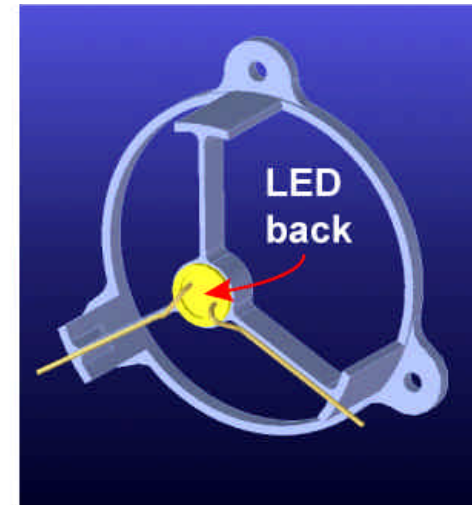
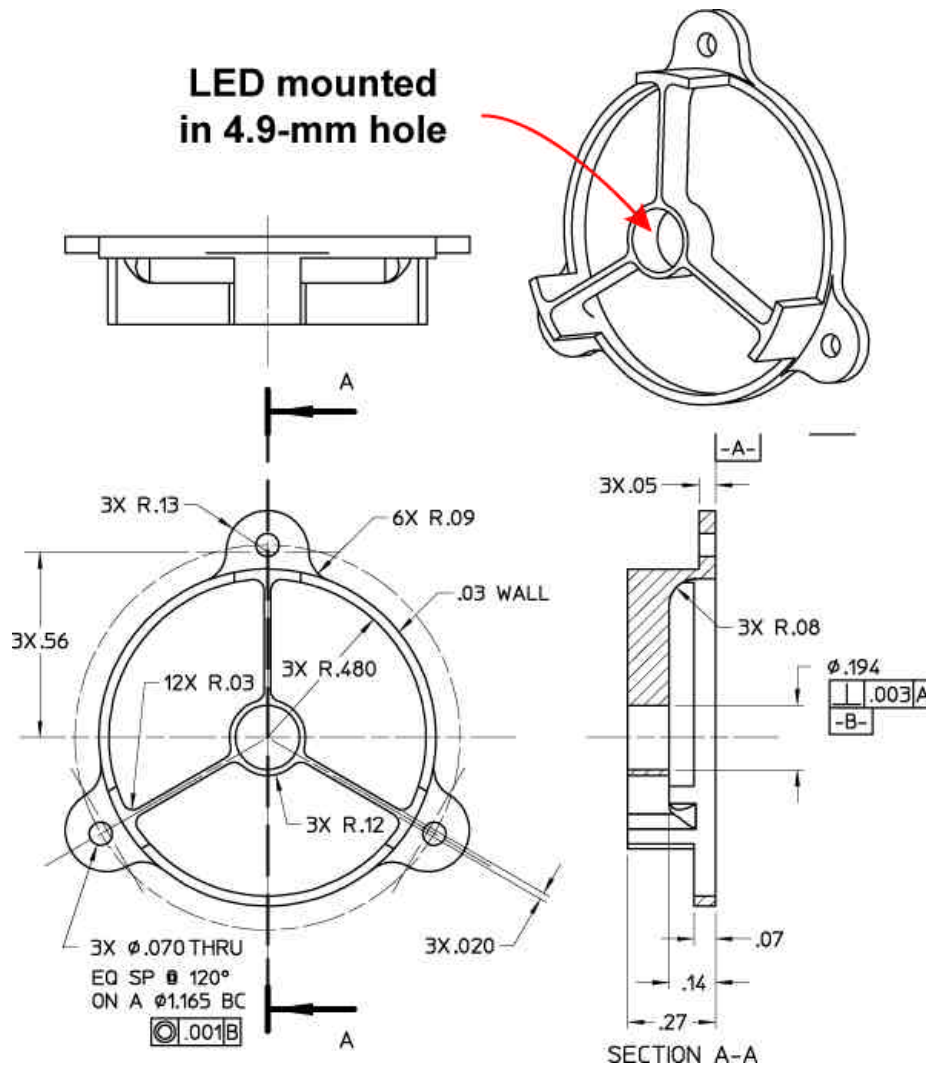


| | |
|--|----------------------------|
| Magnification | 2 |
| Mirror to LED | 15 mm |
| Mirror to image | 30 mm |
| Mirror diameter | 27.51 mm |
| Mirror depth | 5.37 mm |
| Mirror semi-major axis | 22.5 mm |
| Mirror semi-minor axis | 21.21 mm |
| LED mount diameter | 7 mm |
| Source divergence | 55° |
| Source obscuration | 25° |
| Image divergence | 29° |
| Image obscuration | 13° |
| LED light collection efficiency | 67% - 18% = 49% |

Elliptical mirrors made by Opti-Forms, Inc.

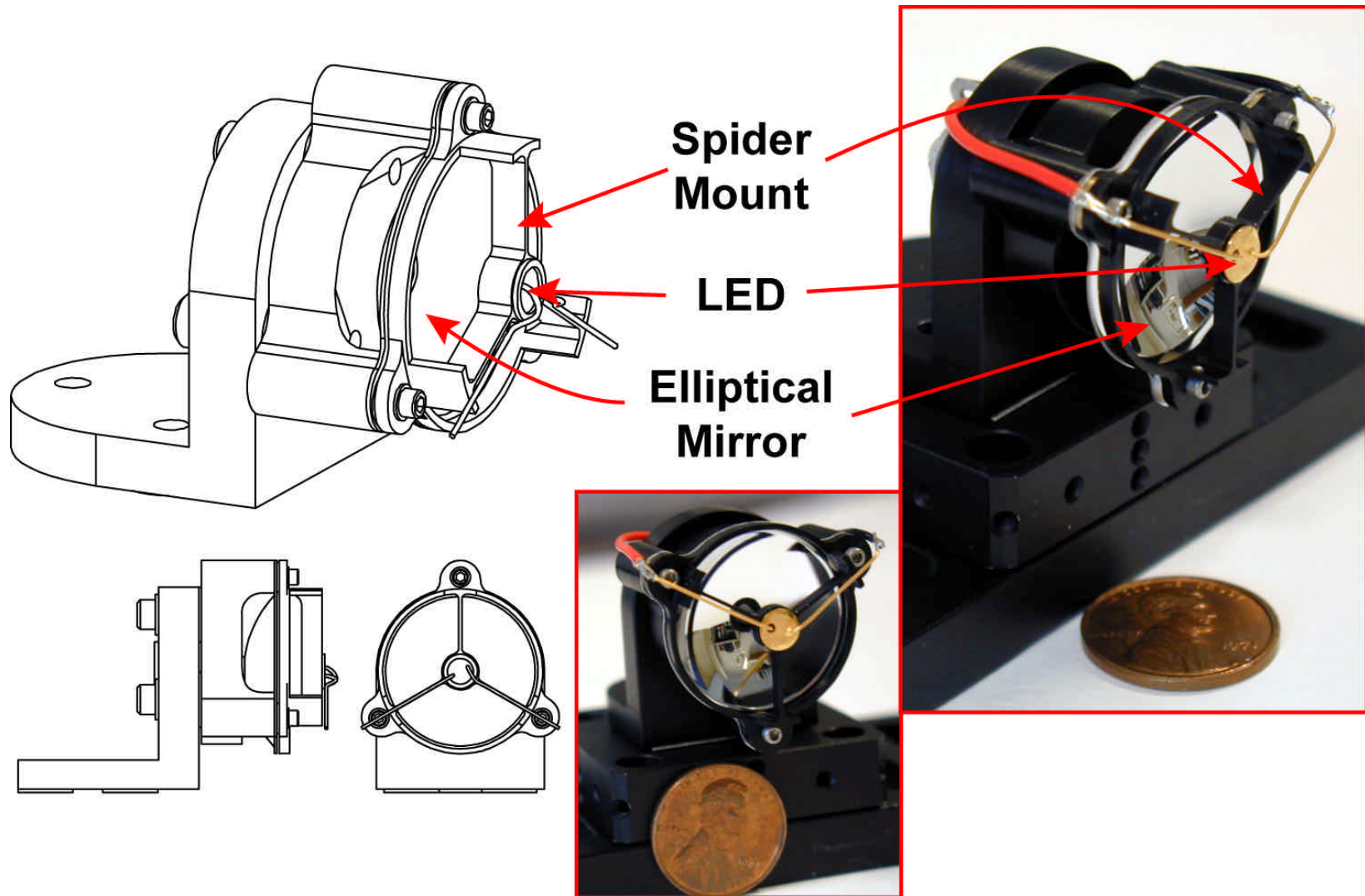


LED "Spider" Mount



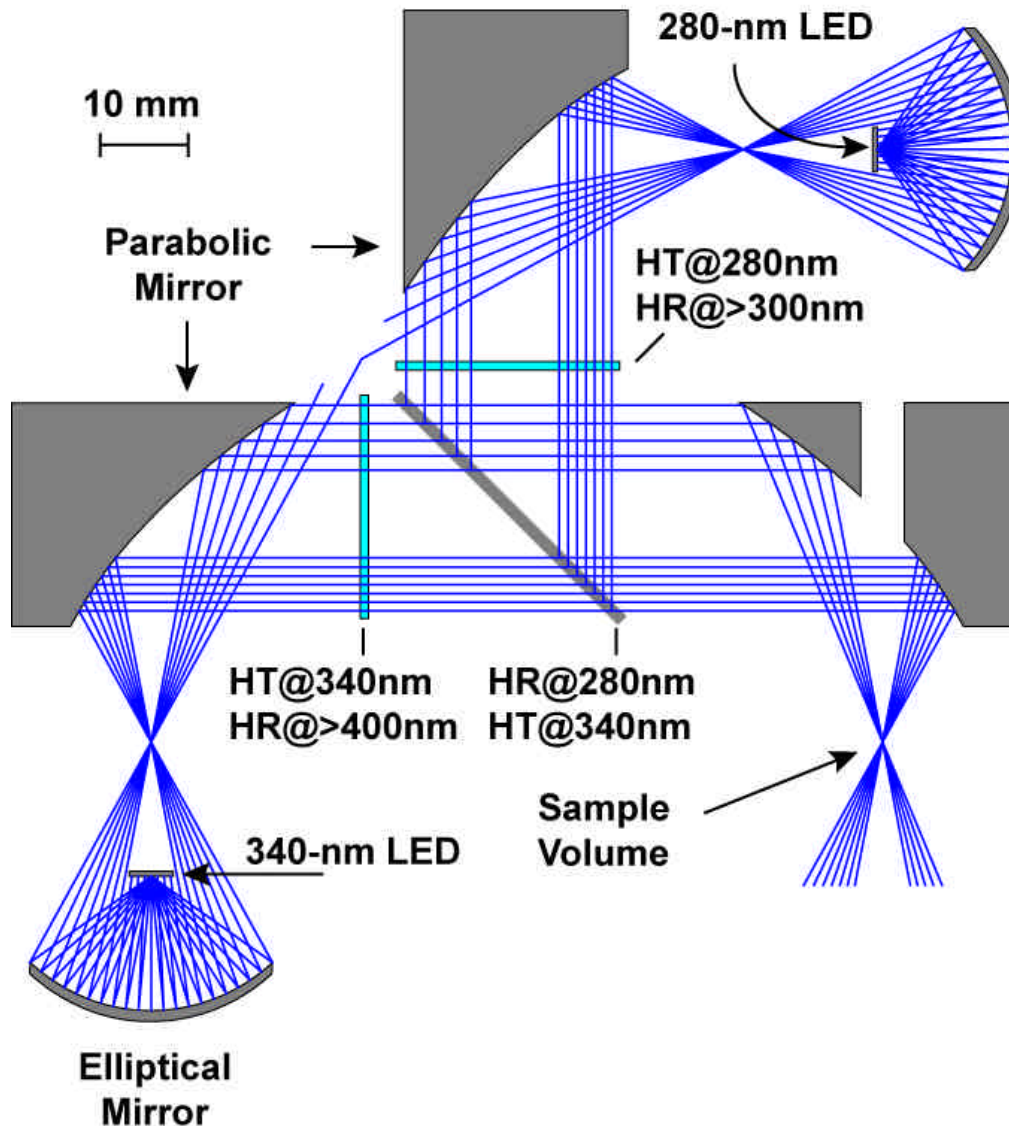


LED Mount and Elliptical Mirror

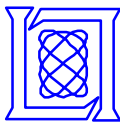




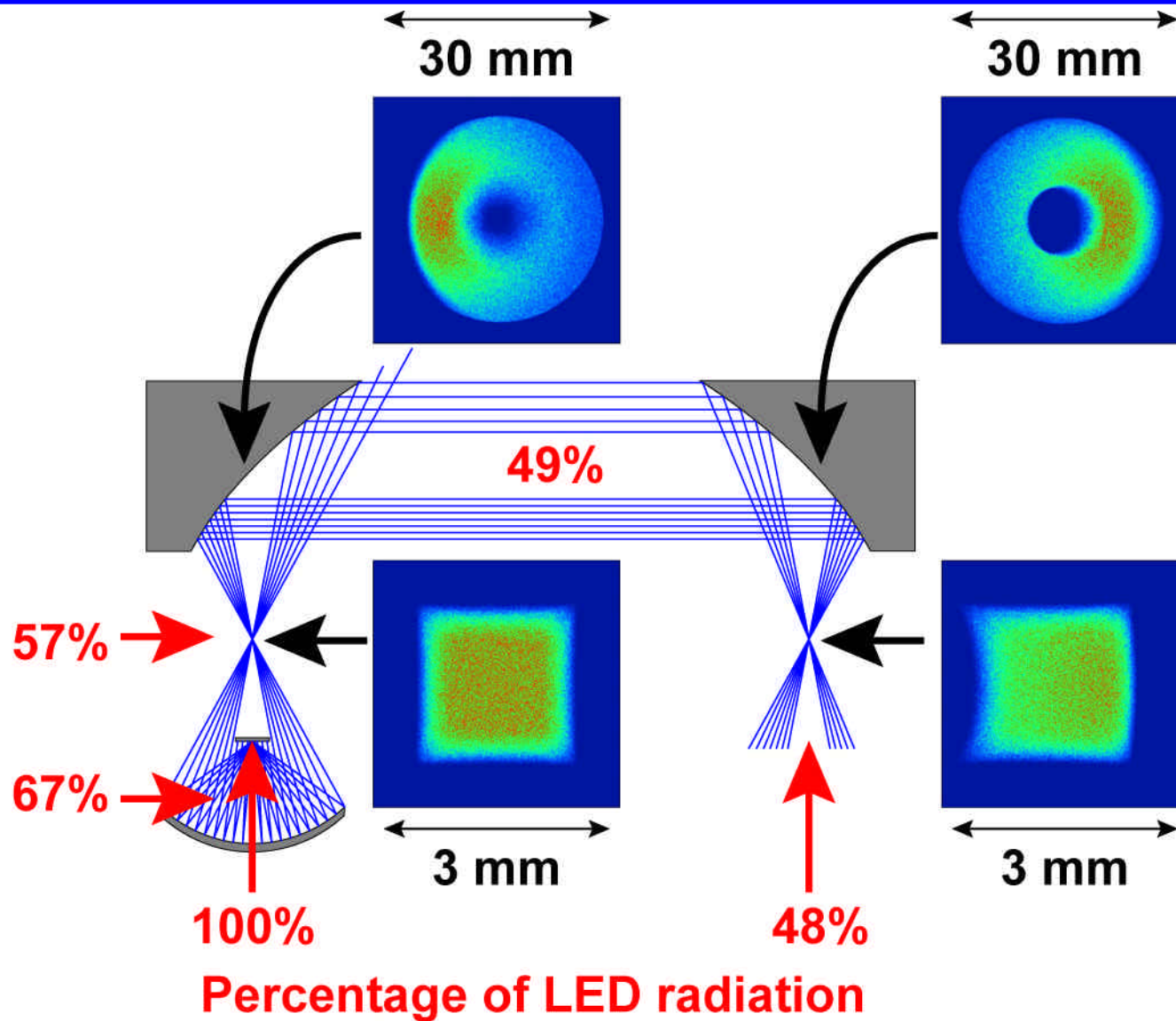
LED Optics



LED diameter = 1 mm
Sample Diameter = 2 mm
Sample Length = 2 mm
Sample Volume = 8 mm³
Transit time = 1.2 ms
Sample Rate = 0.4 l/min

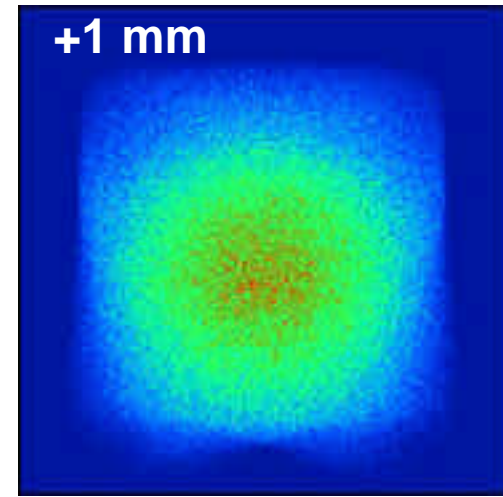
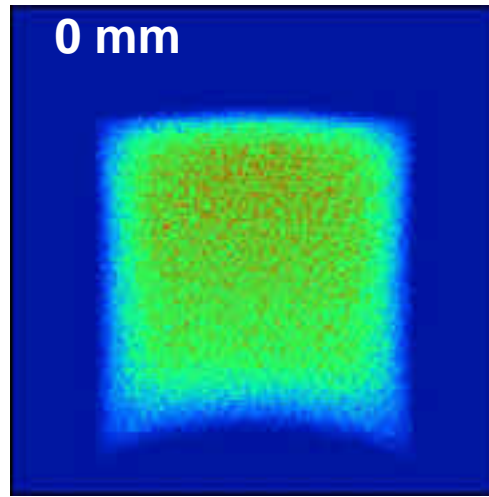
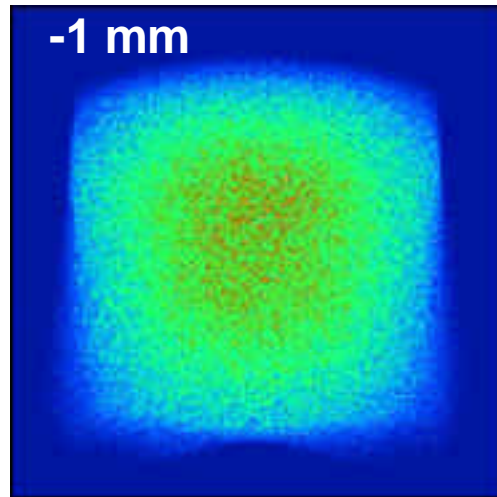


Calculated Imaging of 1-mm LED into Sample Volume

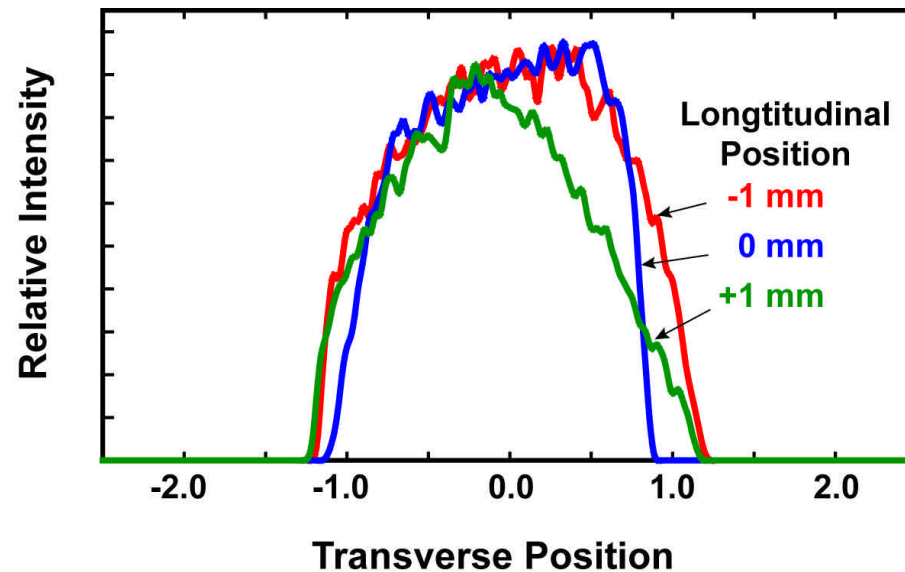




Calculated LED Image Quality in Sample Volume

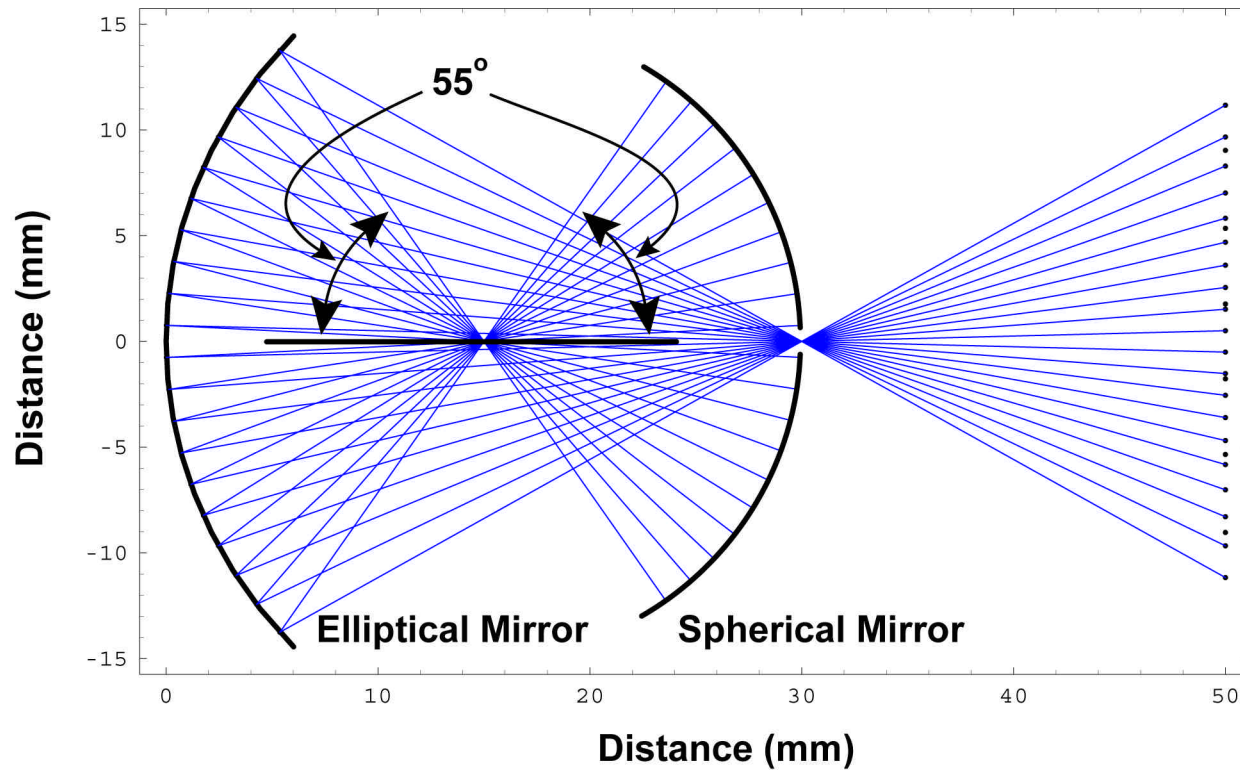


**LED
Source
1 mm x 1 mm**





Signal Collection Mirrors



Collection efficiency = 42%.

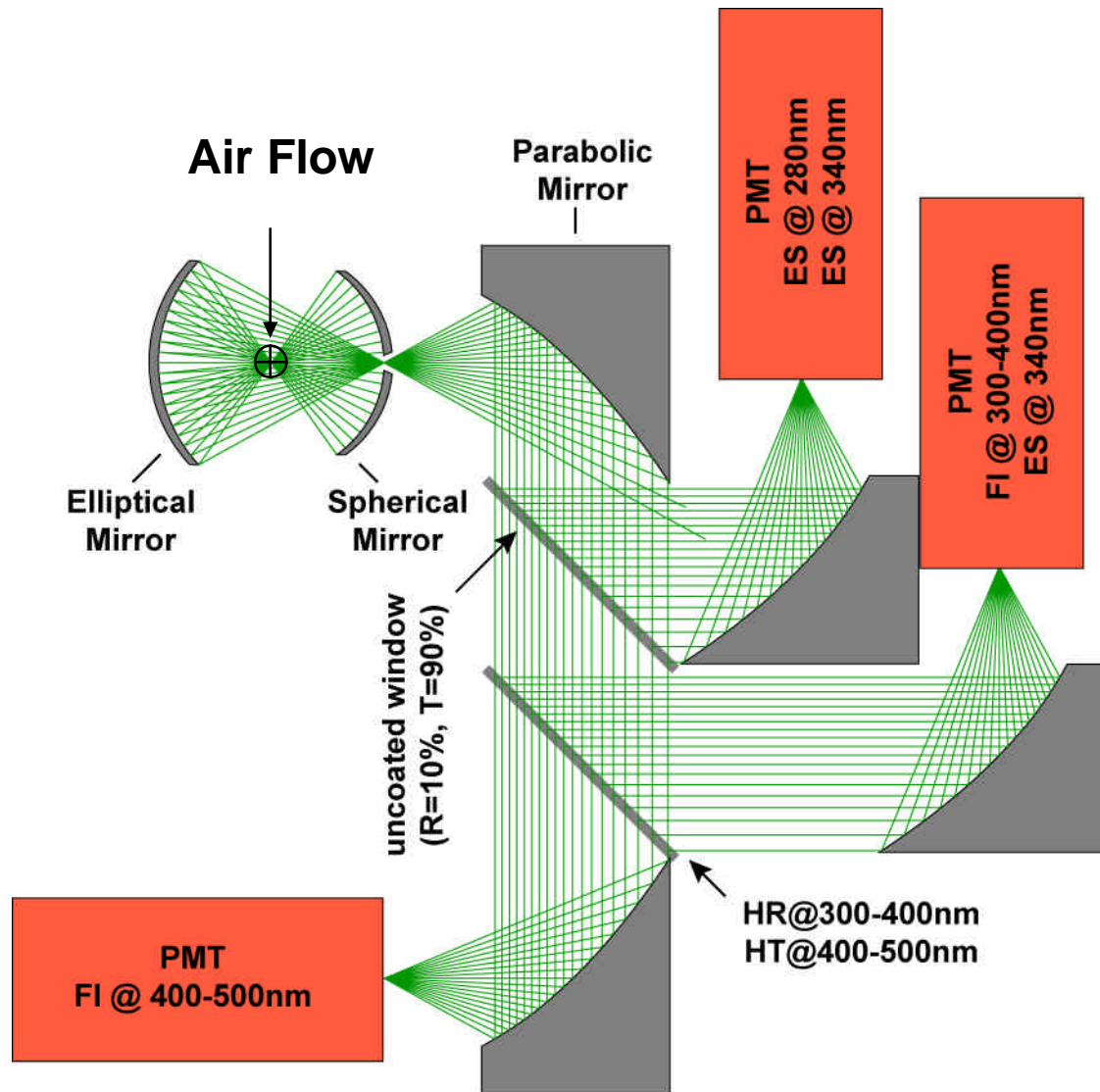
The elliptical mirror is the same as the LED light collection mirror.

The spherical mirror has a radius of curvature of 15 mm.

The spherical mirror has a diameter of 25 mm.

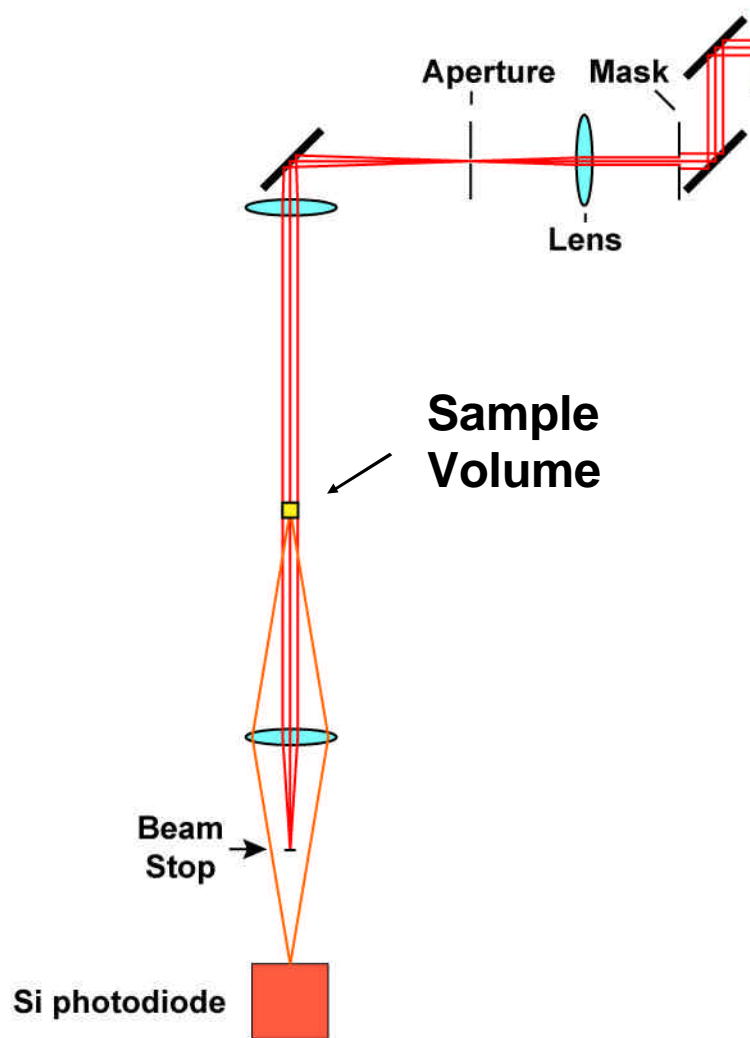


Signal Optics

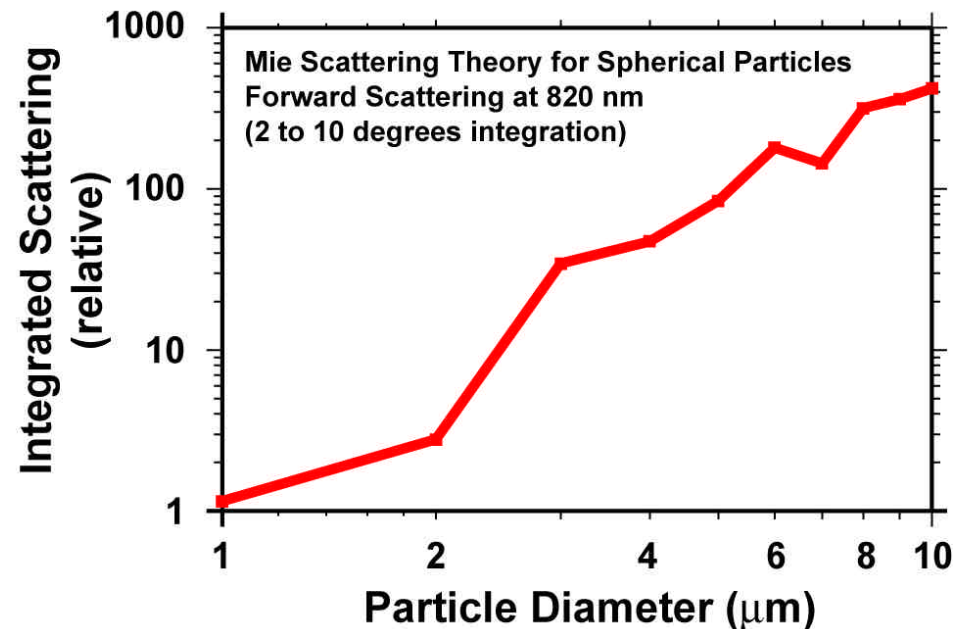




IR Laser Diode

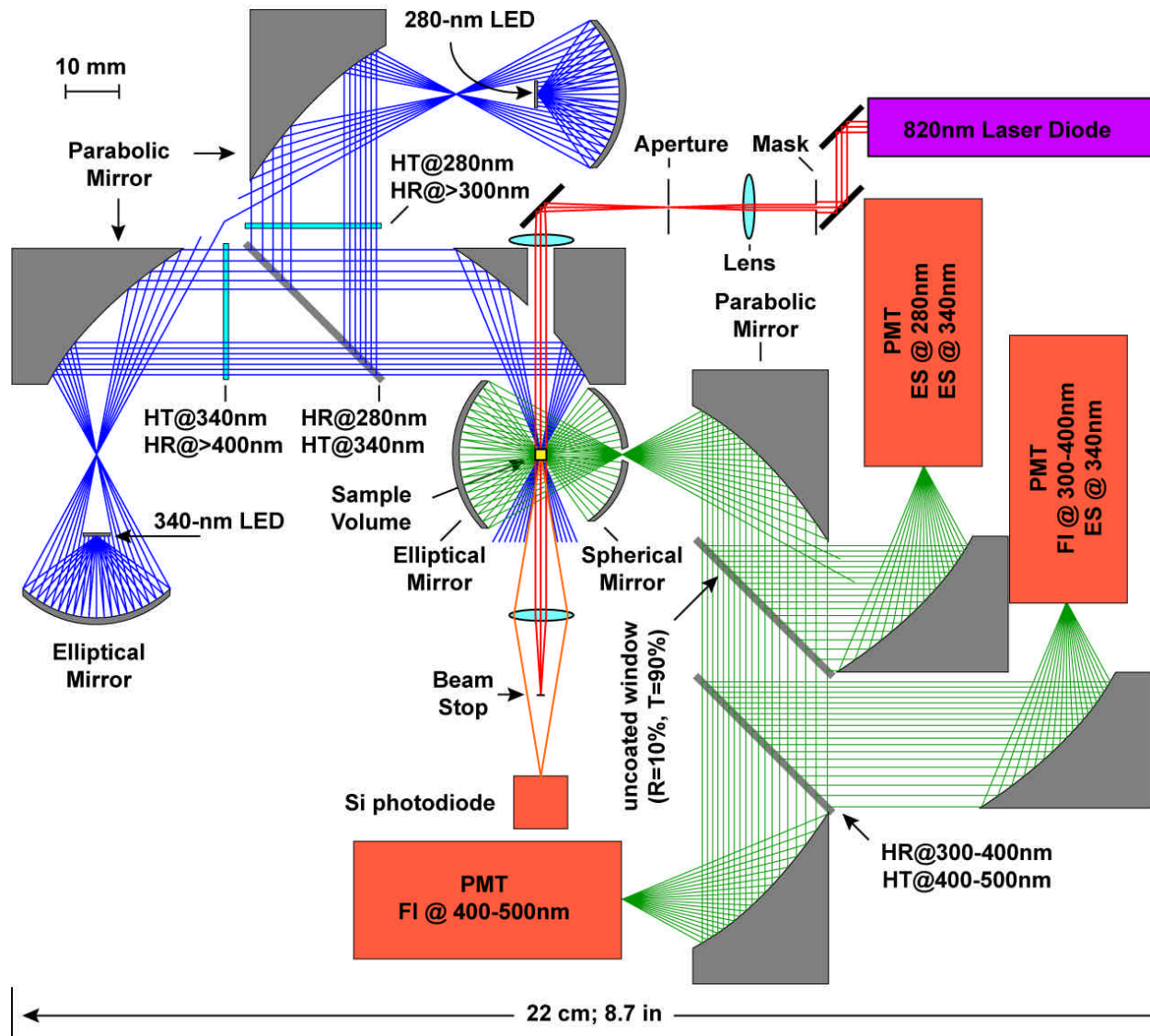


- CW detection of particles
 - Elastic scattering
- Turn on UV LEDs
- Measure particle size
- Measure air speed



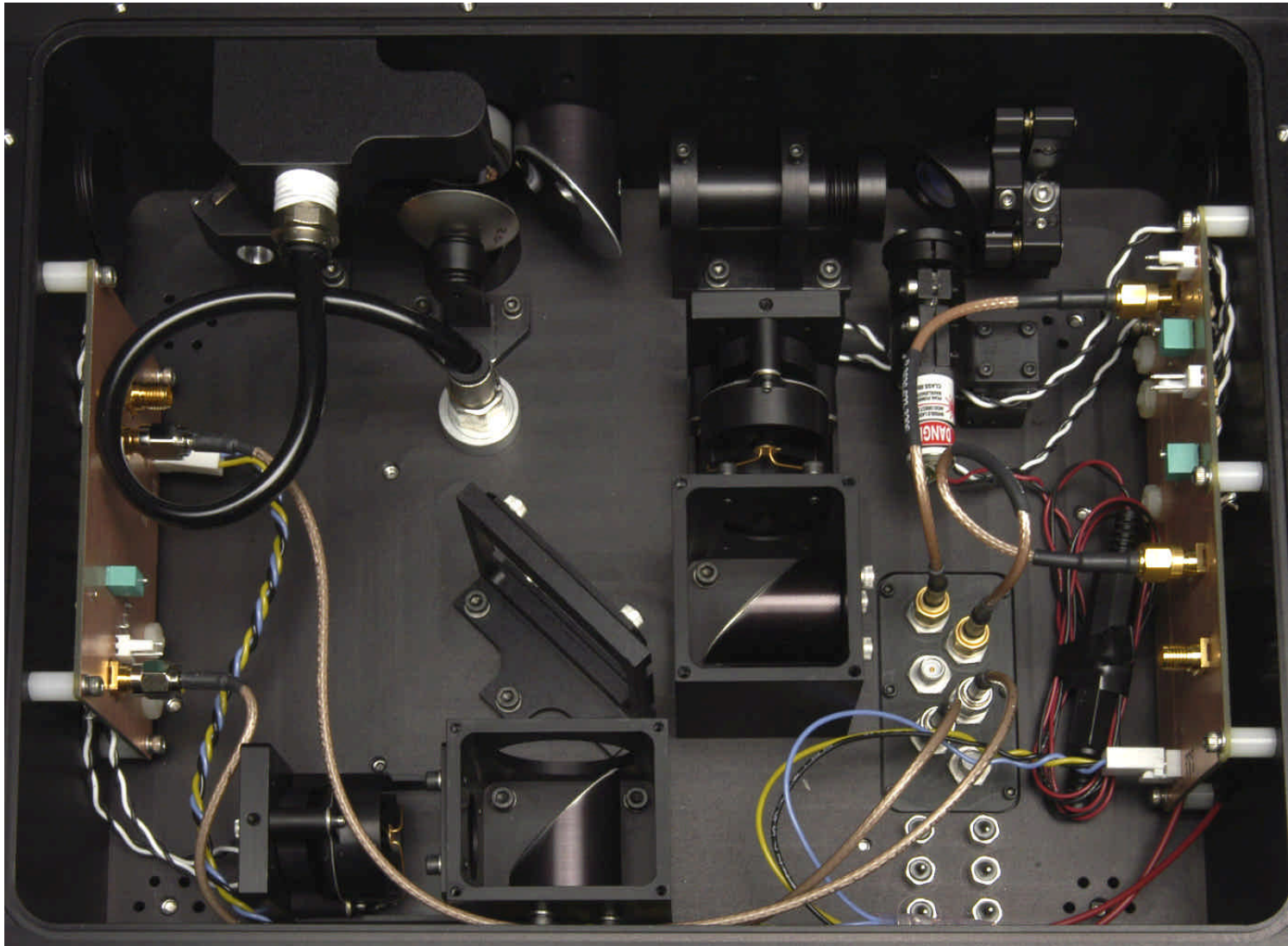


Combined Optics



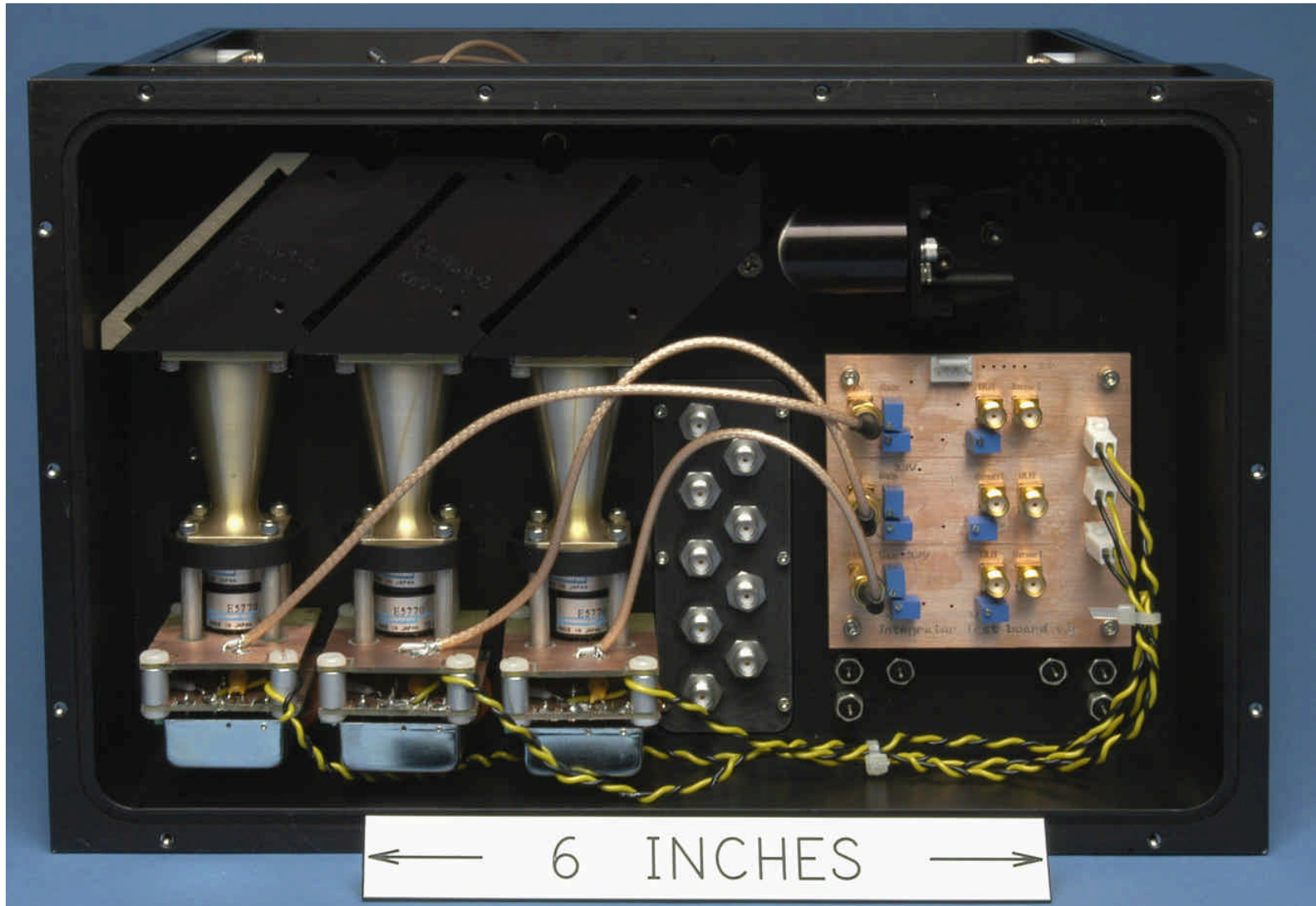


EDU Optics Chamber





EDU PMT Chamber





Summary

- **High efficiency optical system**
 - 50% LED light collection efficiency
 - 40% signal light collection efficiency
- **LEDs mounted on good heat sinks**
- **Optical system has minimal distortion**
 - Retains flat top intensity of LED in sample volume
- **IR forward scattering sizes particles.**
- **Large sample volume**
 - No concentrator needed
- **Compatible with clean-optics air flow design**

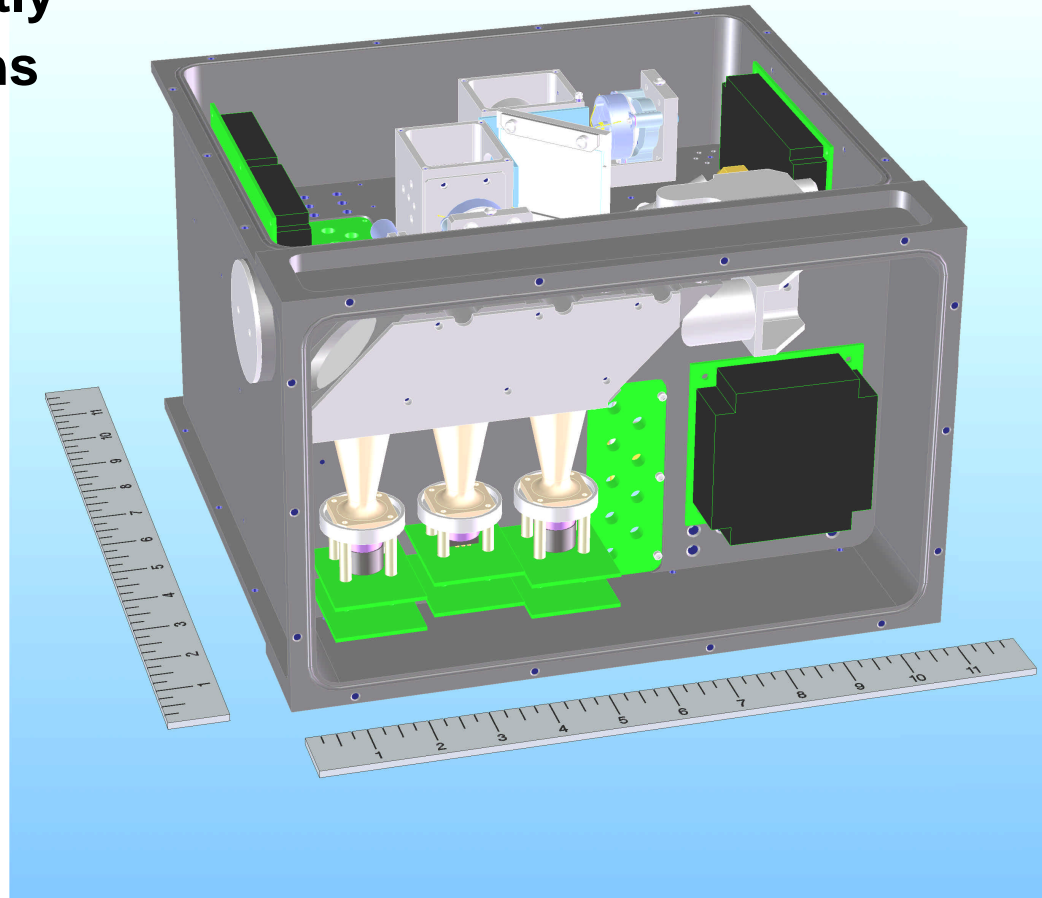


Biological Agent Sensor Testbed Mechanical Design



Mechanical Design

- **Collect LED light efficiently**
- **Collect signal light efficiently**
- **Utilize two LED wavelengths**
 - LED modulation
 - PMT gating
- **Keep optics clean**
 - Confine sample air flow
- **Small size, low weight, low power**
- **Low cost**
 - LEDs
 - Everything else





Mechanical Systems

- **Housing**
- **Airflow System**
 - Requires a non-turbulent stream of air to be sampled
 - It is a “Suck-Thru” system, with control valves on Sample & Make-up air inlets
 - Simple “set and forget” control
 - Filtered make-up air
 - Makeup air delays turbulence of jet
 - Maintains cleanliness of optics
- **IR Triggering system**
 - Forward scatter of IR laser light off particles in sample volume is used to trigger the LED's
 - Extends the life of the LED's
 - Provides particle size information
 - Also triggers the data collection system

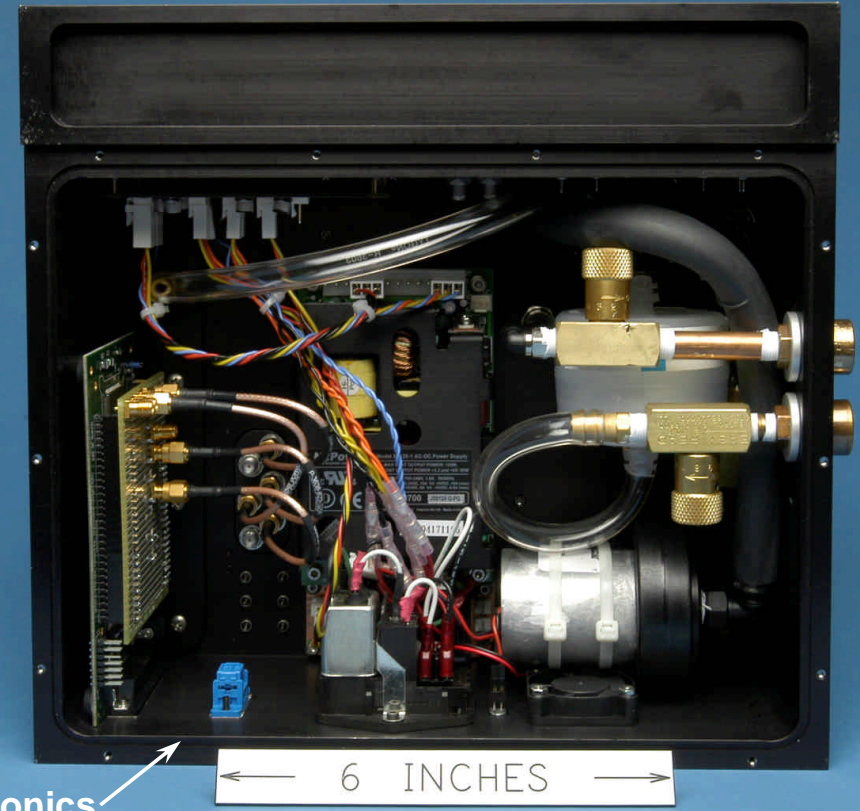
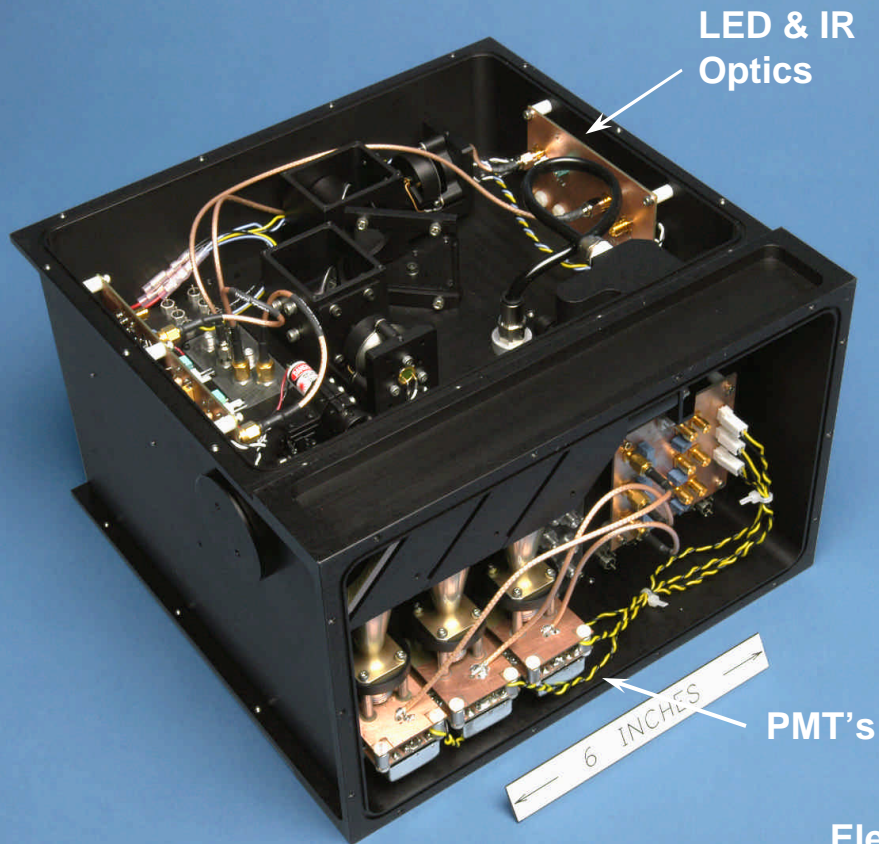


Mechanical Systems con't.

- **LED light sources to illuminate particles**
 - 340 nm and 280 nm light sources to alternately illuminate the sample volume
 - Light sources are optically co-aligned and filtered while collimated
 - Designed optical mounts to be as self aligning as possible for a EDU
- **Scattered and Emitted light collection**
 - Light from particles in the sample volume is collimated and then wavelength separated to three PMT's
 - Elimination of stray light from LED sources is extremely important



SUVOS/BAST Housing

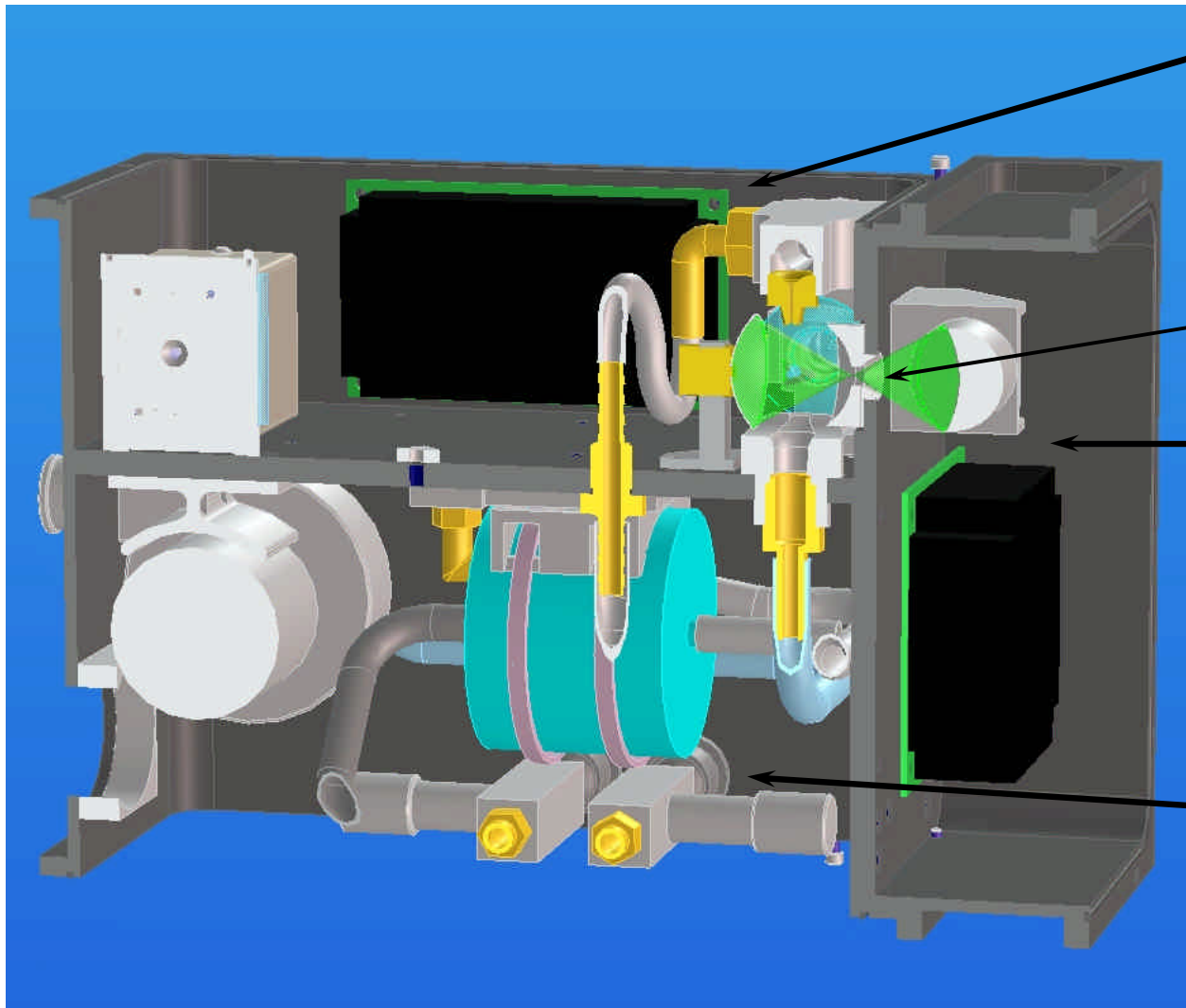


Designed to be compact
Divided into three compartments
Provides easy access to all compartments

Weight approx. 17.5 lbs
11.1" x 10.9" x 7.3"
Can be used in any orientation



Cross Section Through Housing



Upper Optical Compartment – Houses LED's and IR Scatter Detector

- Light-tight
- Airtight (except for Inlets & Outlet)
- Maintains cleanliness of optics
- Only open to PMT Compartment at Sample Chamber Opening

PMT Compartment

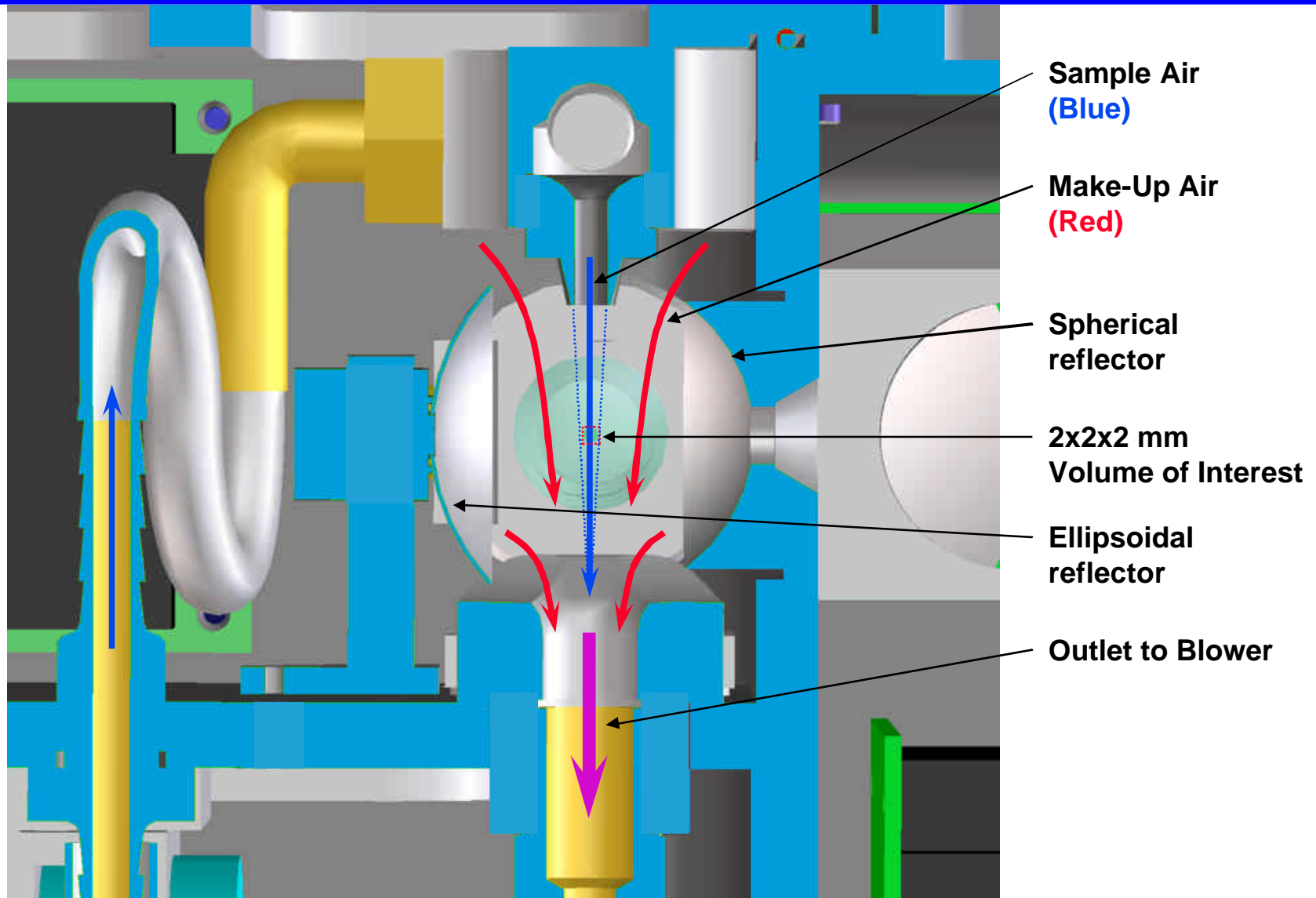
- Light-tight
- Airtight
- PMT's are away from stray light
- Keeps high voltage in one area

**Electronics Compartment
Contains Blower & Ducting**

- Fan Cooled



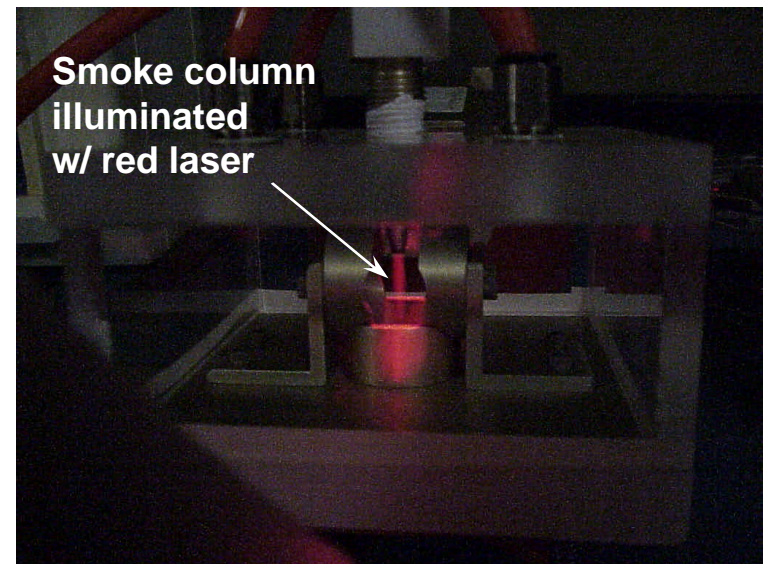
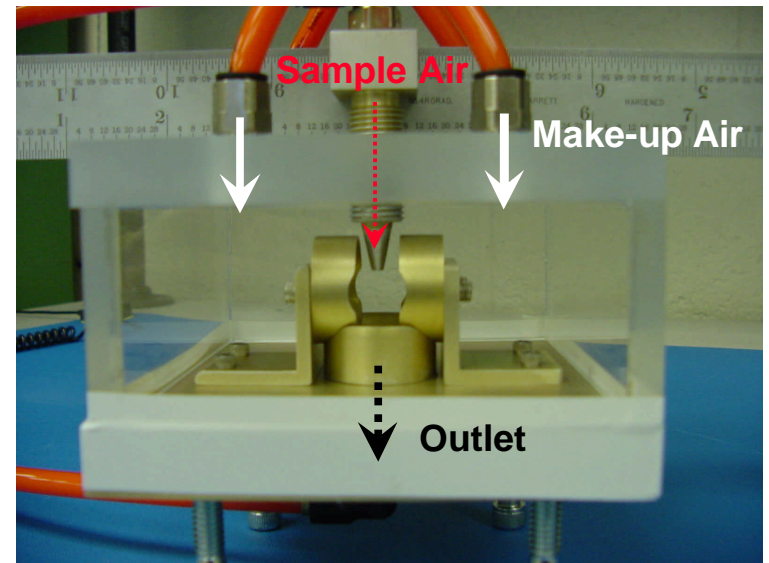
Cross Section Through Sample Chamber





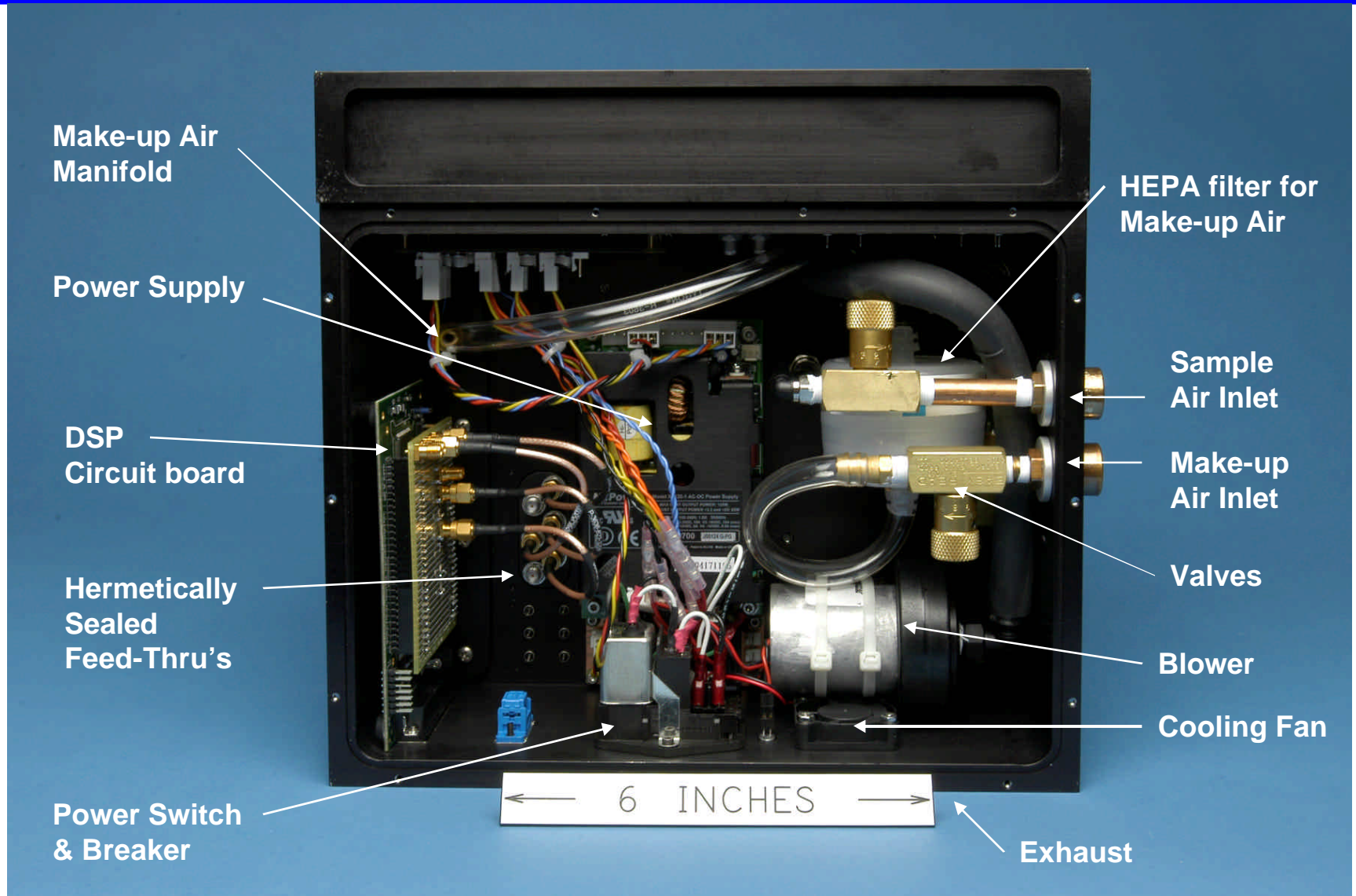
Test Chamber to Verify Airflow

- **Require sample air jet to be laminar**
 - Provides a straight particle trajectory
 - Prevents dirty optics which would cause scattered light
 - Allows accurate measurement of airflow velocity
- **Test model was used to help analyze flow**
 - Design uses a inlet nozzle with make-up air supplied from outside the collector optics
 - Smoke was used to allow visualization of airflow
- **Make-up air flow rate is 10X the sample air flow rate**



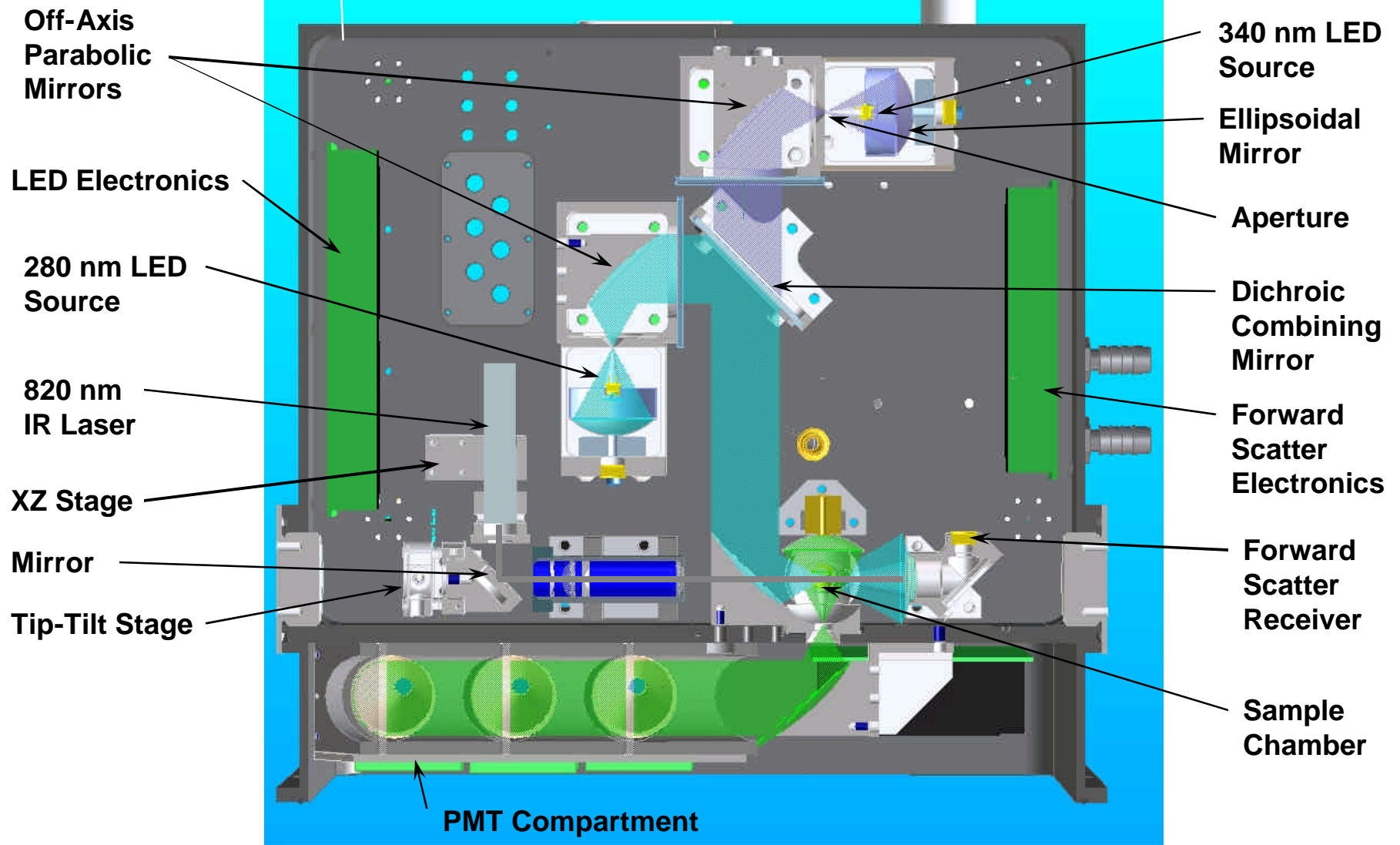


Electronics Compartment



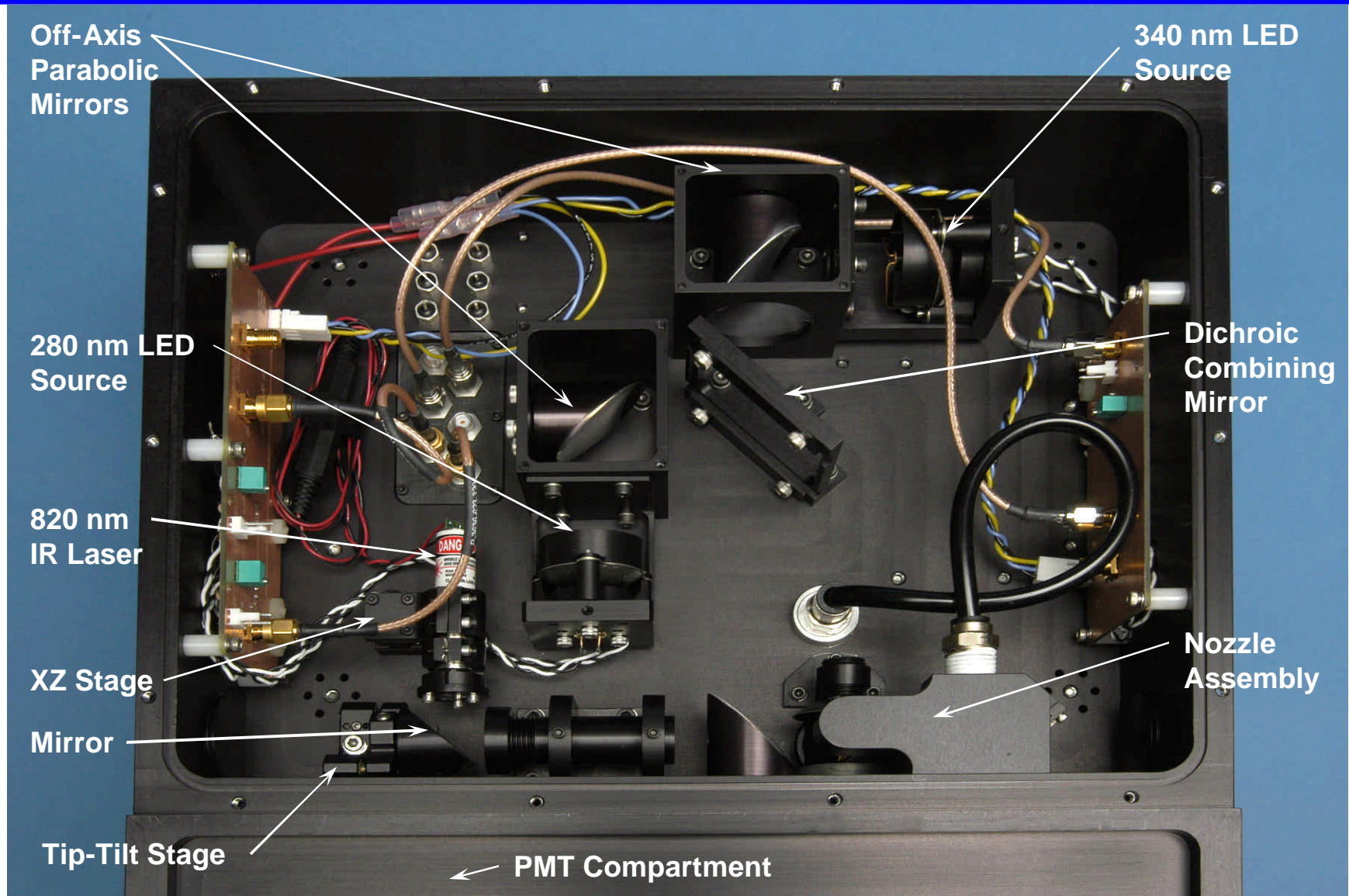


Section showing Mechanical Layout of Optics



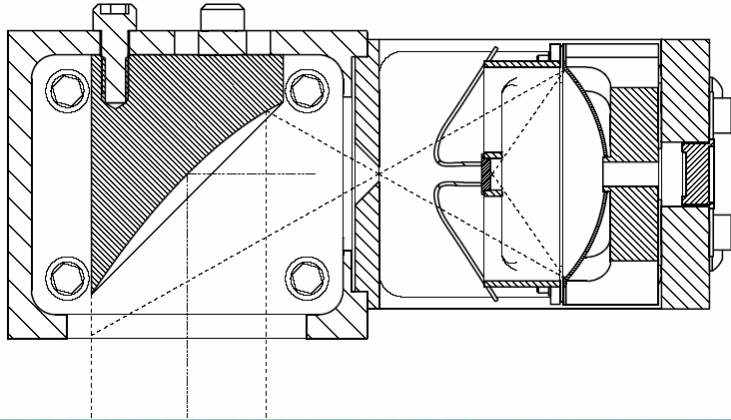


Optics Compartment



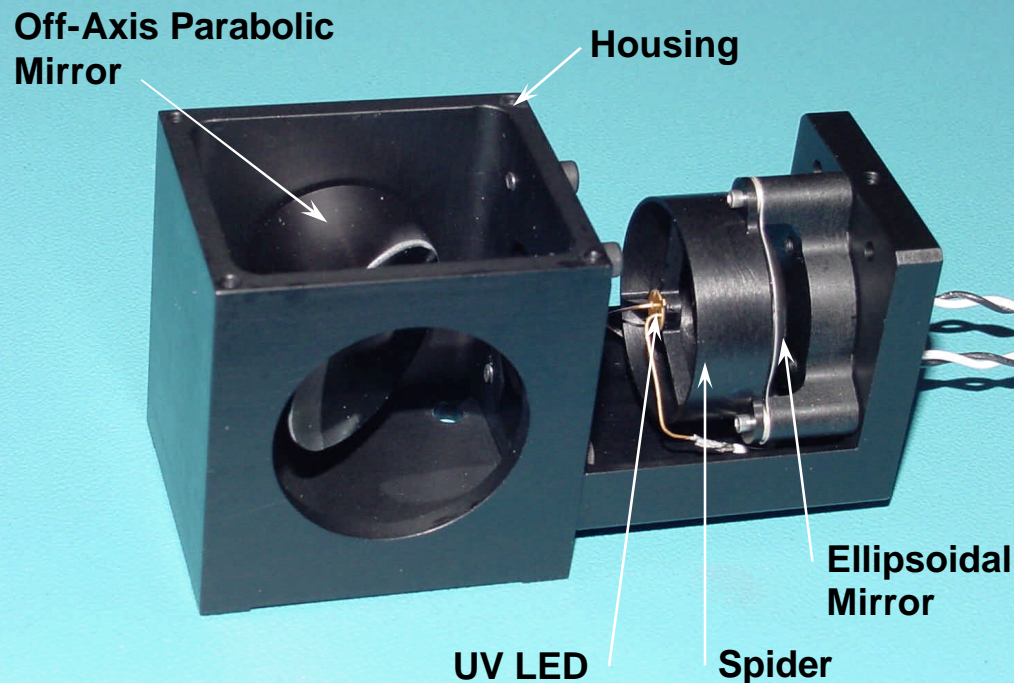


Detail of UV LED Source



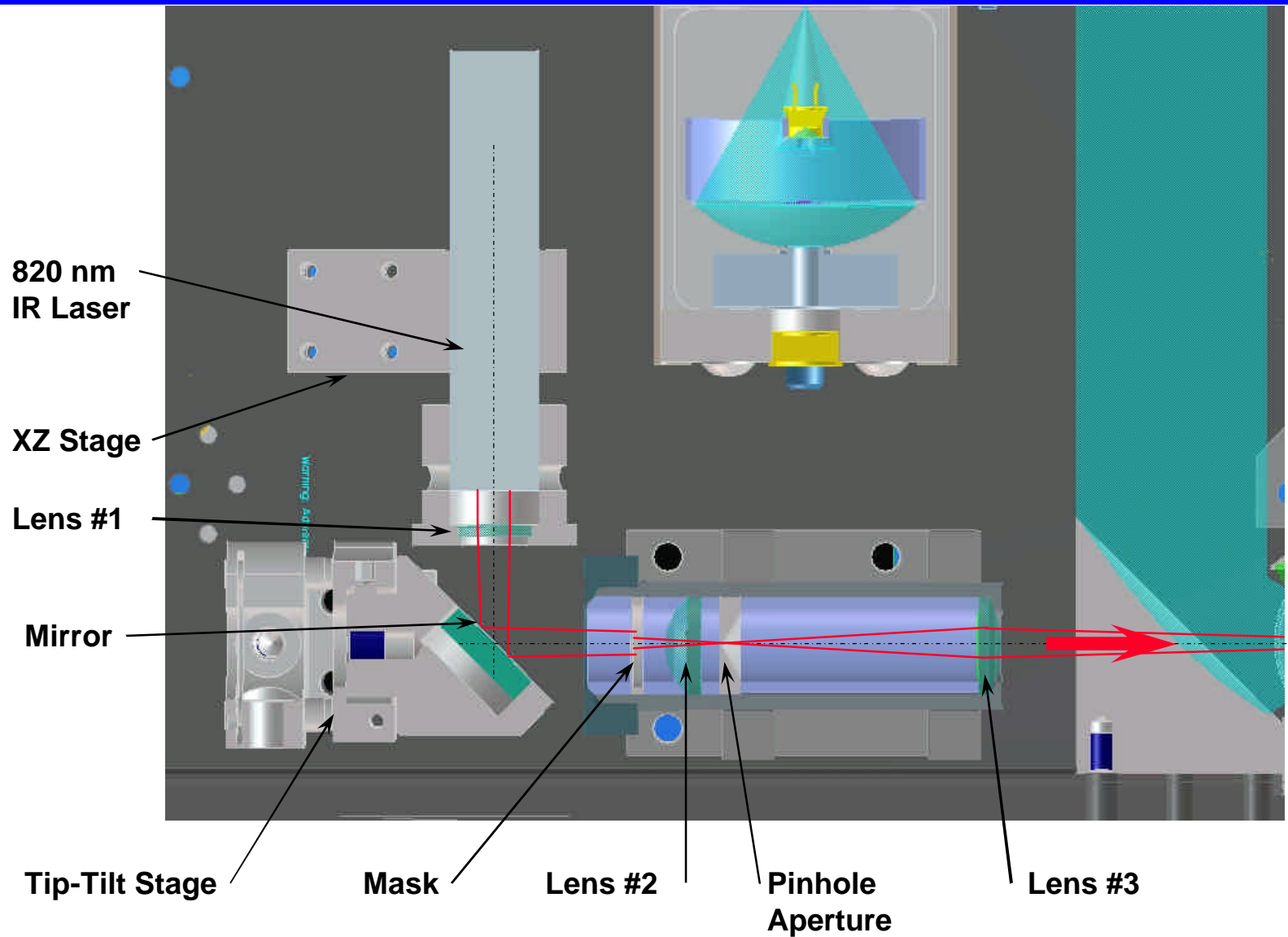
LED Source design

- Large collection angle
- Easy to align and focus
- Interchangeable mounting
- Eliminate stray light
- Allow monitoring of LED



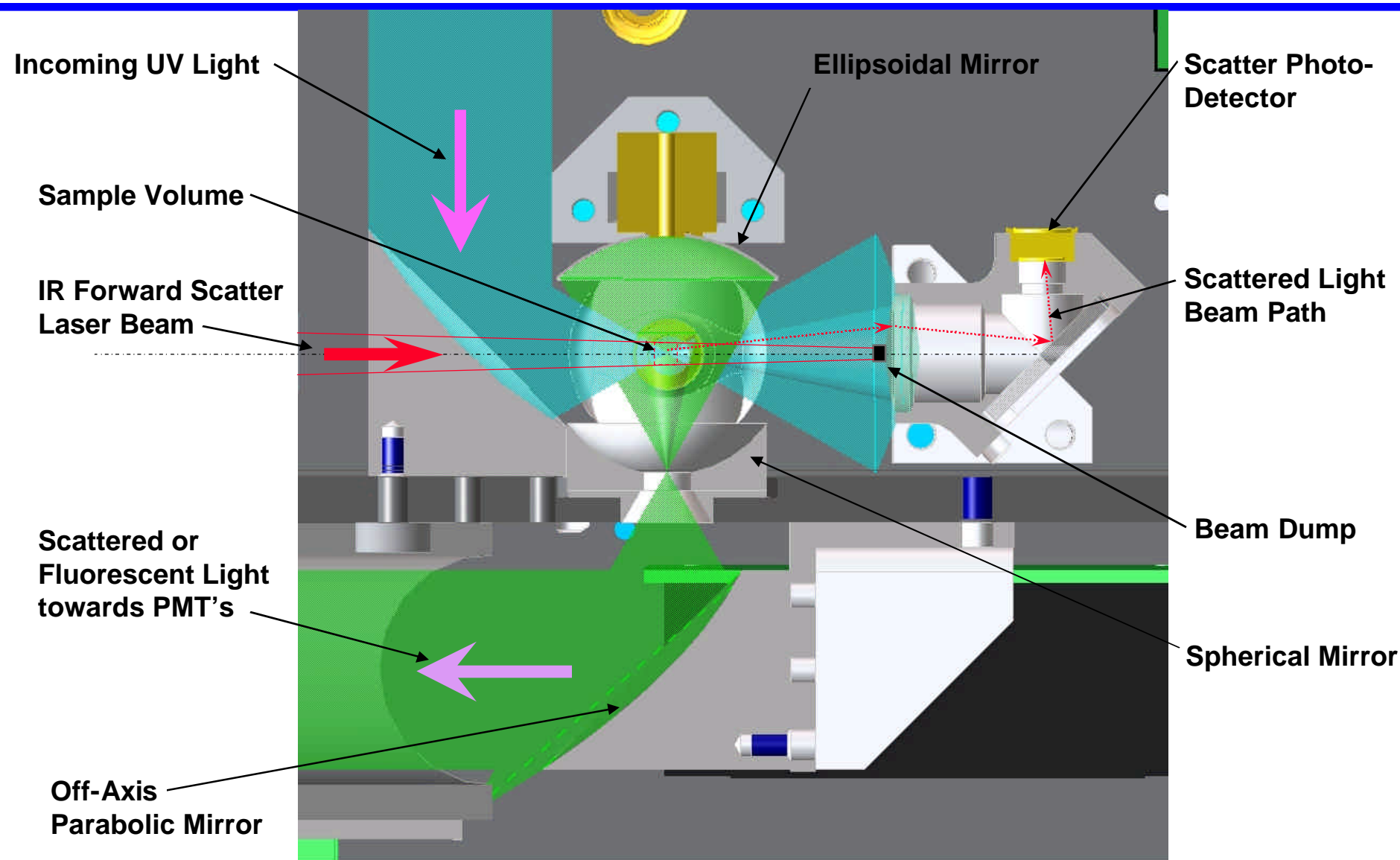


IR Forward Scatter Laser Beam Forming



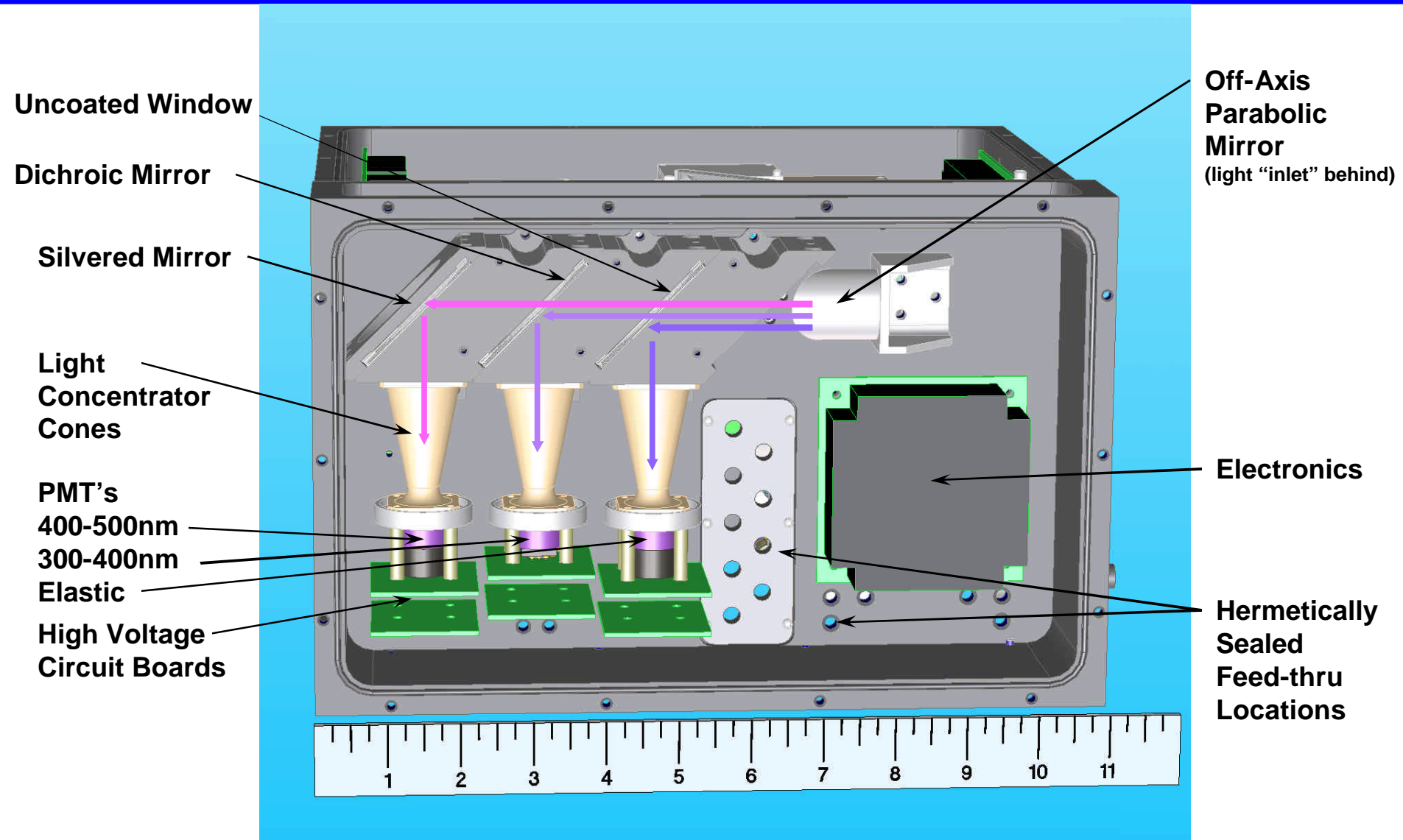


Sample Chamber Optics



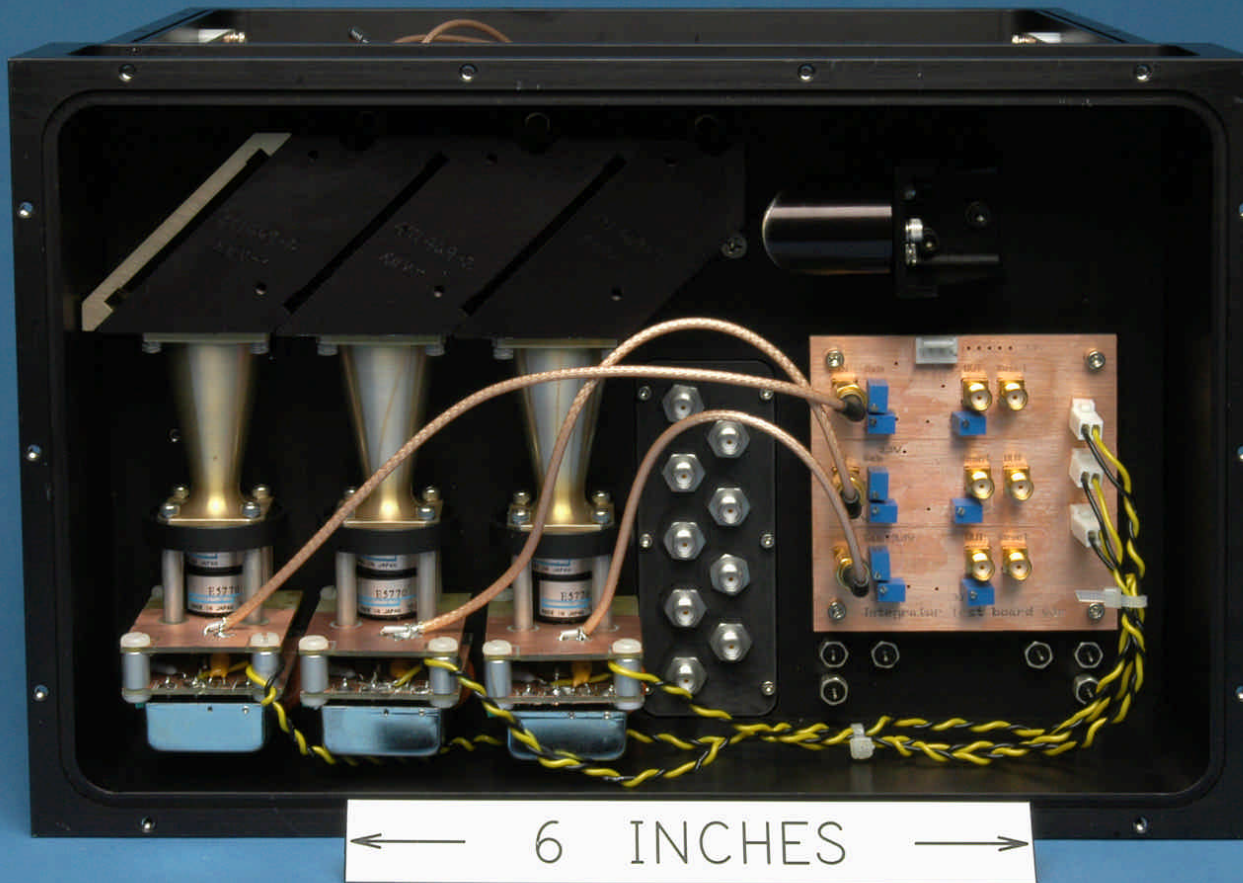


PMT Compartment Layout



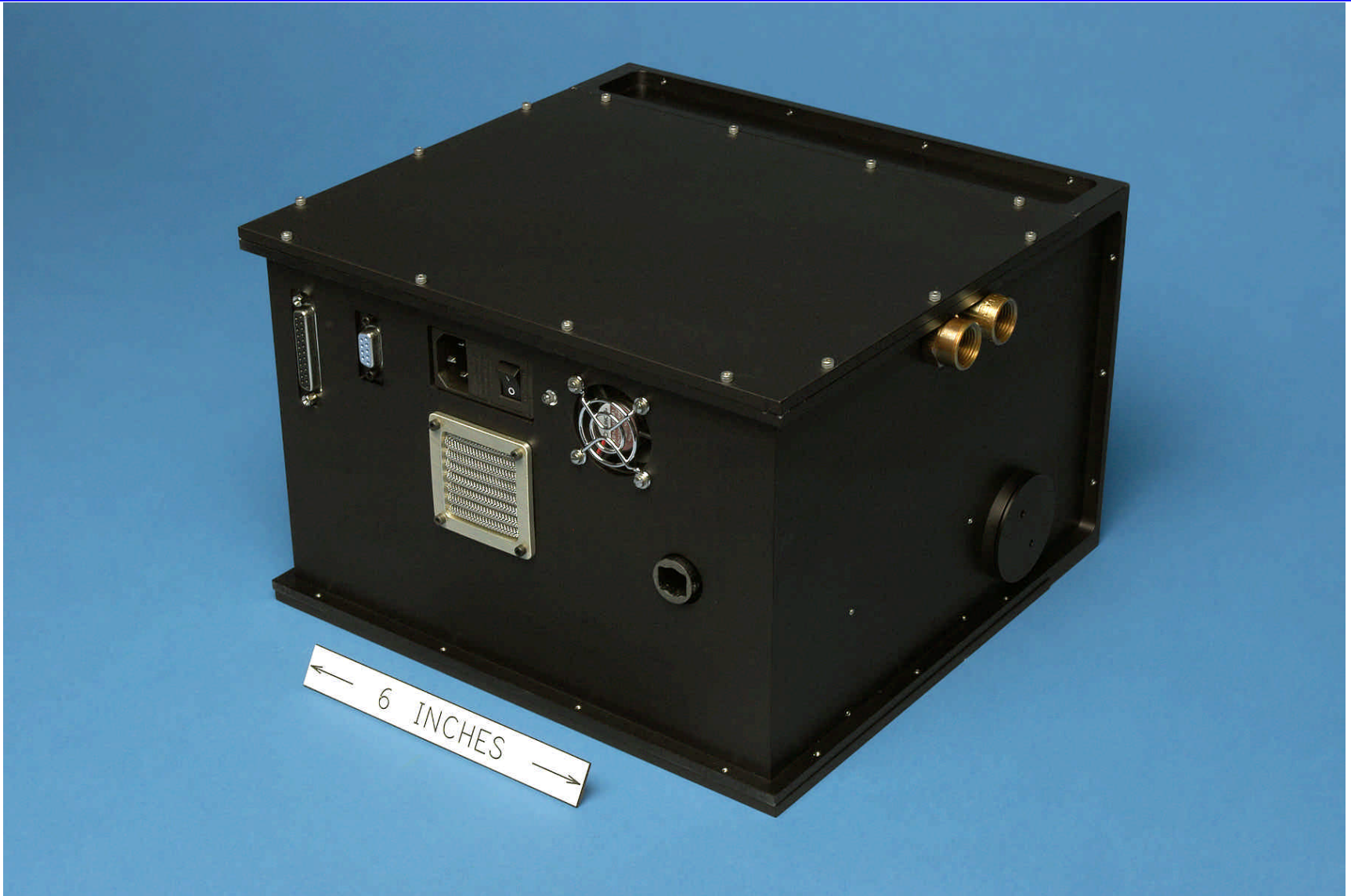


PMT Compartment





SUVOS/BAST with Covers Installed



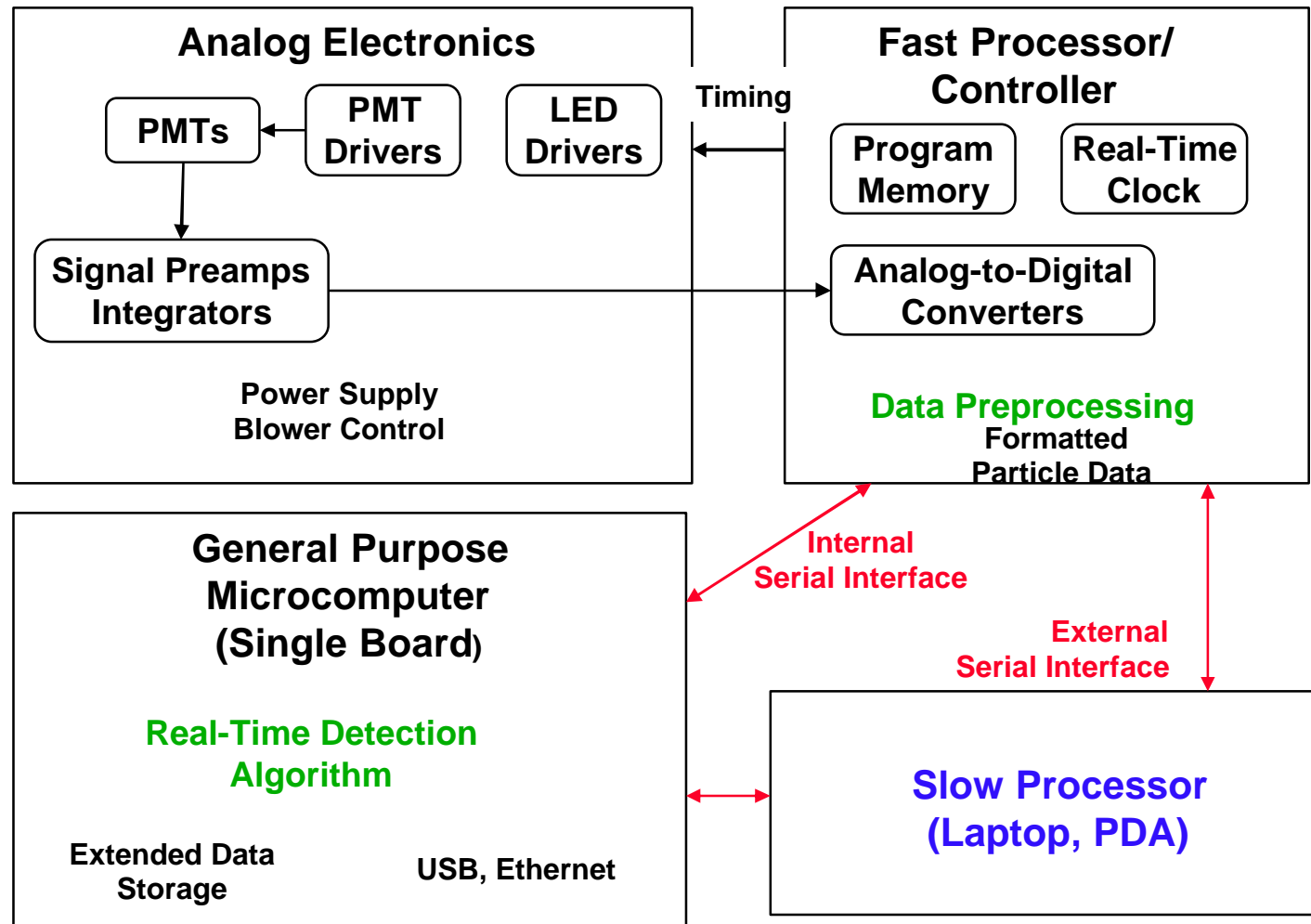


Biological Agent Sensor Testbed Electrical Design



Engineering Development Unit Design: Electronics

Electronics Block Diagram





Outline

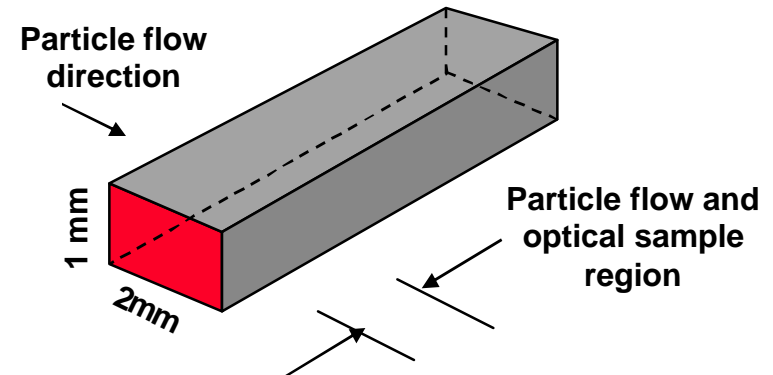
- **System operation**
 - Trigger and UV beams
 - Signal capture
- **Hardware design**
 - Design philosophy
 - Block diagrams
 - Board layouts
- **Summary**



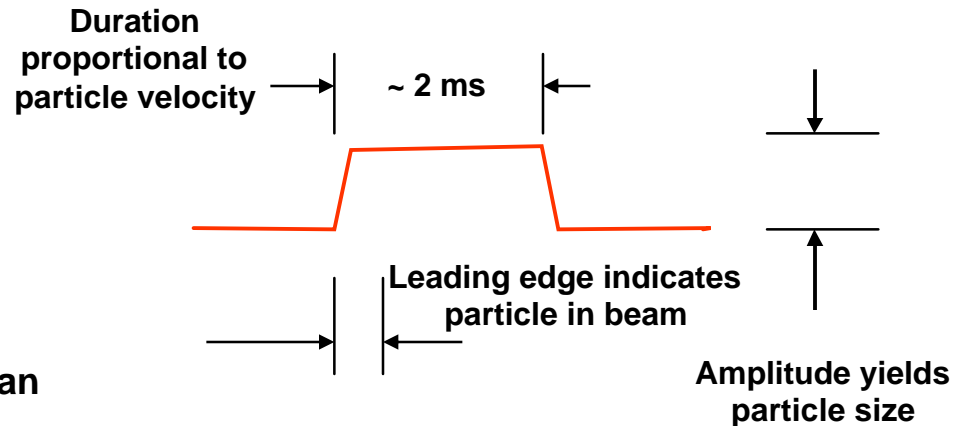
Trigger Beam

- **Excitation**
 - 820 nm CW laser diode, 8 mW
 - Conditioning optics creates flat top beam profile
- **Detection**
 - Forward elastic scatter 2 to 10 degrees
 - Silicon photodiode with integral amplifier
- **Sampling rate**
 - 60 kHz
- **Functions**
 - Particle detect
 - Trigger UV sources
 - Particle size estimate
 - Primary particle descriptor
 - Velocity
 - Yields correction factor for time in beam
 - Average of many particles can be used to control blower

Beam Geometry



Output Signal





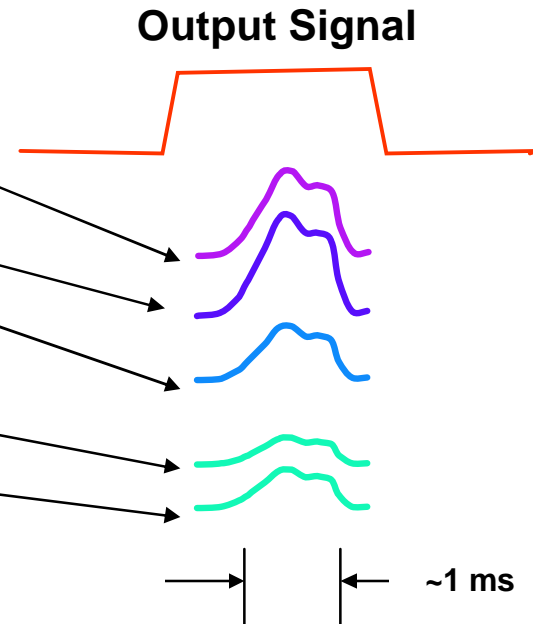
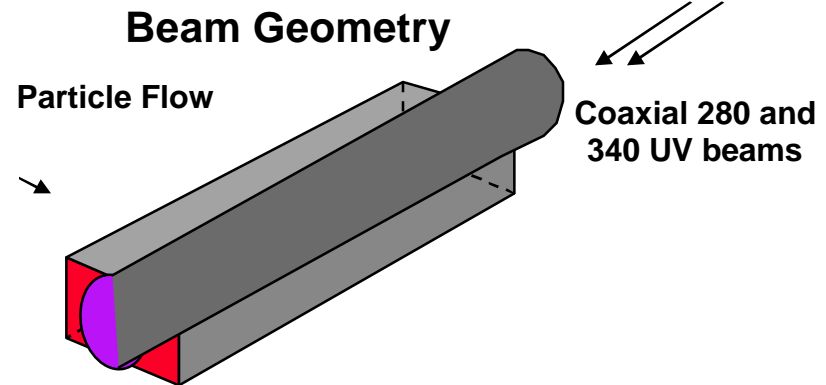
UV Beams

- **Excitation**

- **280 nm LED**
0.15 mW CW, 0.45 mW pulsed
- **340 nm LED**
0.8 mW CW, 8 mW pulsed

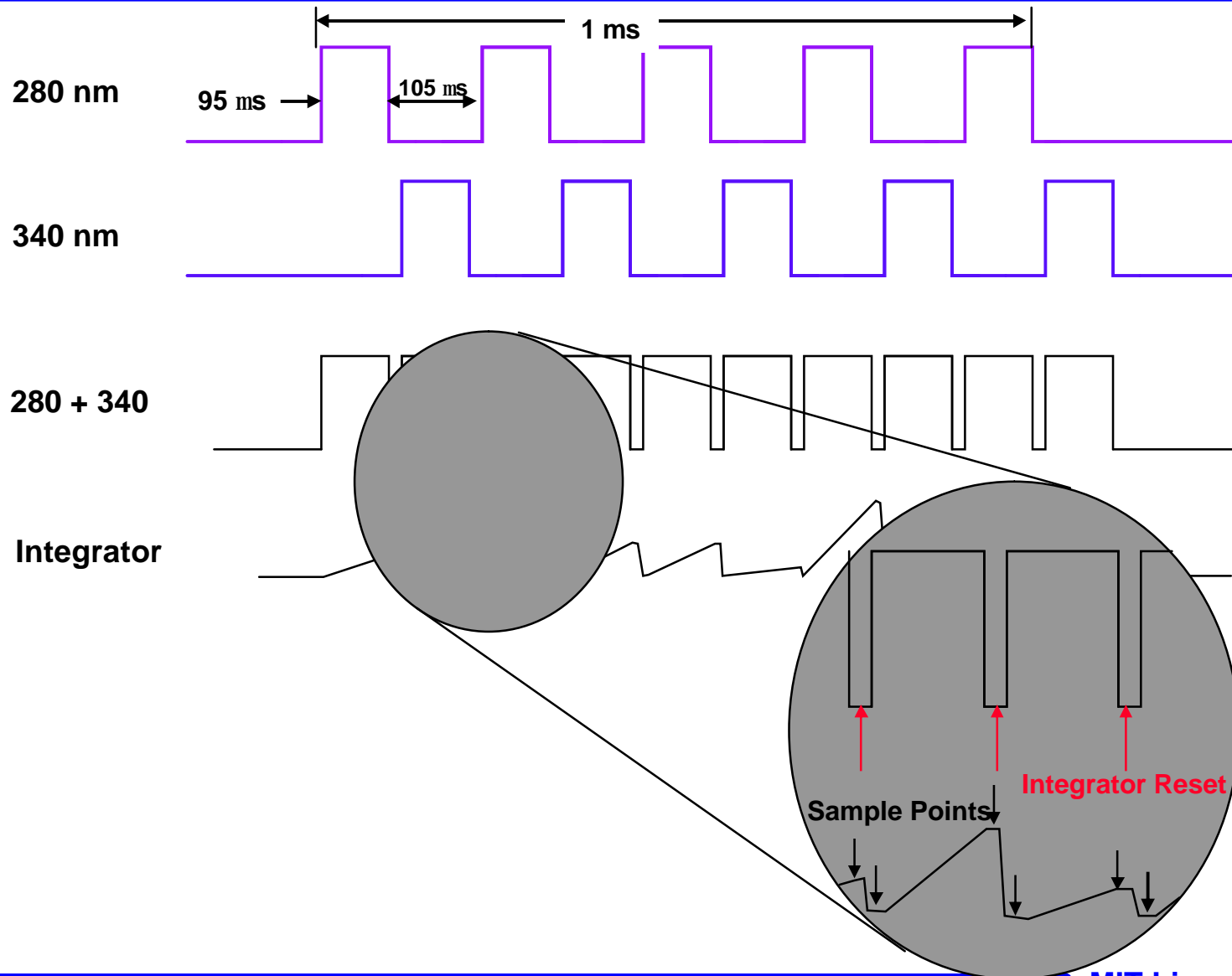
- **Detection**

- **PMT 1, 280 nm**
Elastic scatter @ 280nm
- **PMT 2, 300 - 400 nm**
Elastic scatter @ 340
Fluorescence due to 280
excitation
- **PMT 3, 400- 500 nm**
Fluorescence due to 280
excitation
Fluorescence due to 340
excitation





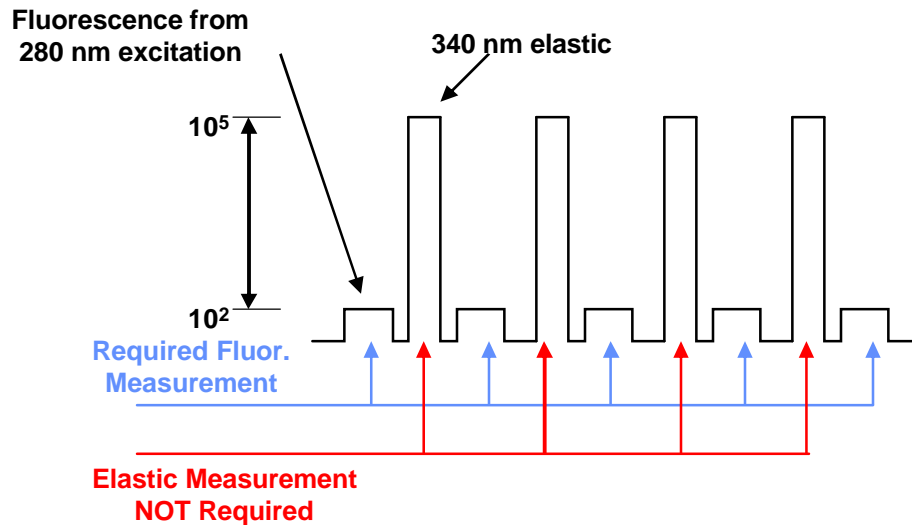
Time-Multiplexed UV Signals



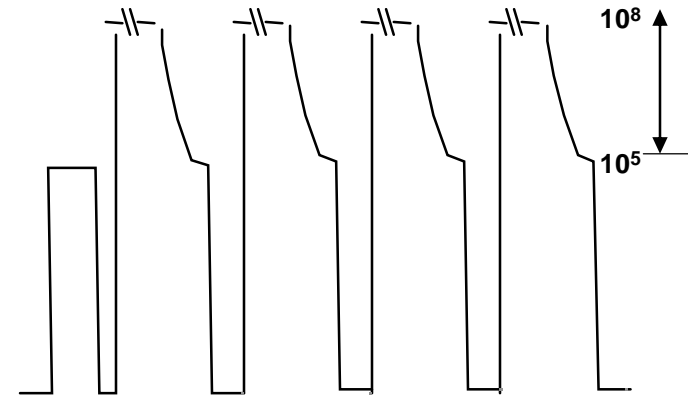


PMT 2 – Saturation Problem

1 mm particle



10 mm particle



- Elastic signal is $>1000\times$ as bright as fluorescent
- PMT saturates on large particles, losing capability to detect fluorescent signature
- Solution – Squelch the gain on PMT when elastic signal is present



Signal Estimation

- **All particle features (size, fluorescence, color) are measured relative to standard calibration particles**
 - Typical cal target: 2 μ m polystyrene spheres
 - Calibration particle features are assumed consistent from particle to particle
 - Removes need for absolute measurements
- **Calibration Phase**
 - Perform instrumented aerosol chamber tests with cal targets
 - Measure conversion factor (C)
$$\text{PPL}_{\text{aerosol}} = C * (\text{Particle counts/sec}), \text{ where } C \text{ is a function of airflow through instrument as estimated by particle velocity } E[V]$$
 - Determine measurement noise using cal target statistics
- **Measurement phase**
 - Correct raw scatter measurements by
 - Particle velocity (time in beam)
 - Intensity of light sources (as measured by LED monitoring circuit)



Outline

- **System operation**
 - Trigger and UV beams
 - Signal capture
- **Hardware design**
 - Design philosophy
 - Block diagrams
 - Board layouts
- **Summary**

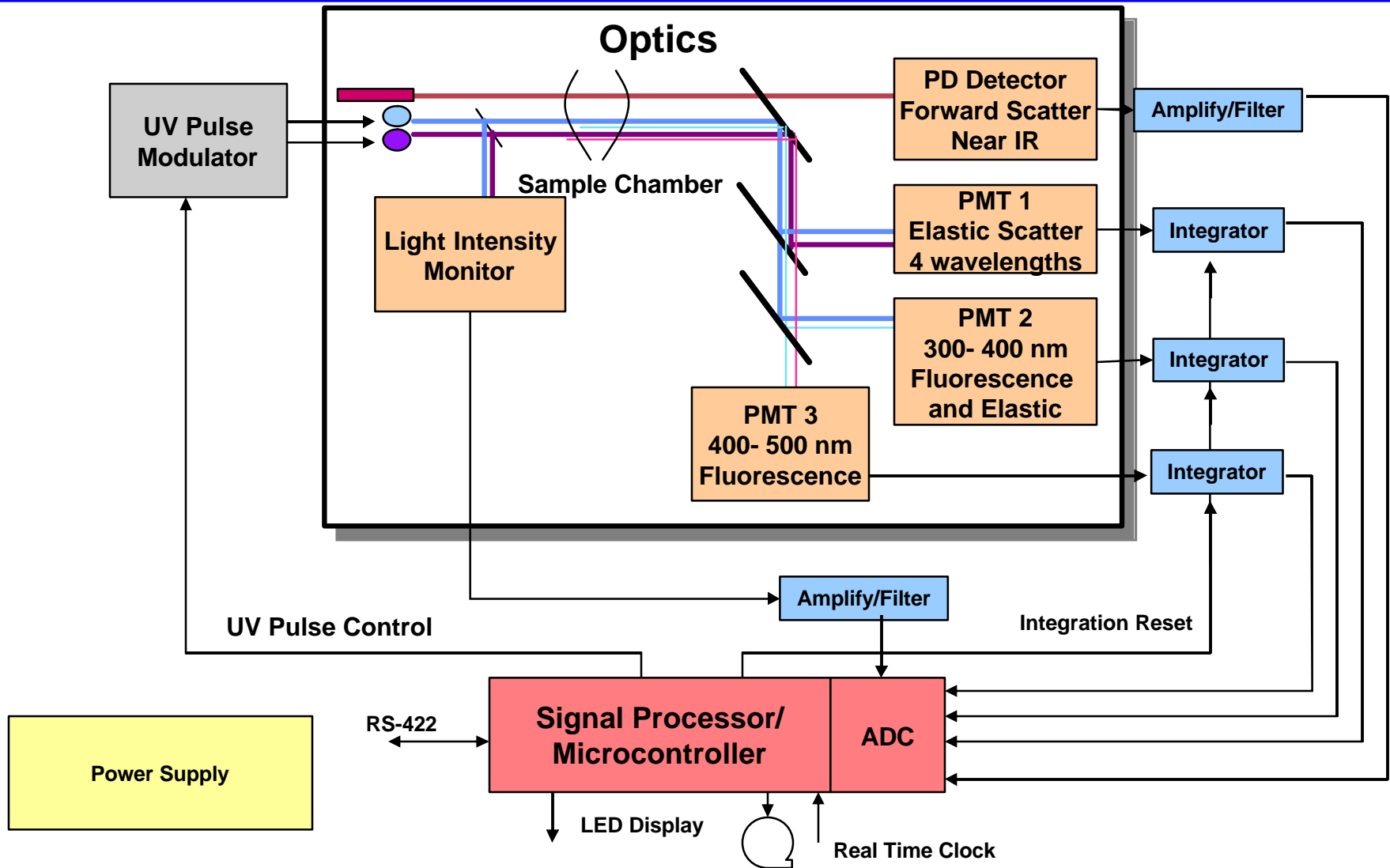


Design Philosophy

- **Use powerful COTS Digital Signal Processor/Microcontroller**
 - Built in timing and processing for time critical decisions
 - Analog-to-digital converters built into circuit
 - Ability to observe any part of code or signals through development workstation
- **“Hand build” simple, distributed analog boards**
 - Single layer boards, routed on prototype machine
 - Small investment in each board - change as needed
- **Data products are vector of characteristics for each particle**
 - Formatted and serially transmit, readable by any general purpose computer
 - Algorithm development and deep memory resides on general purpose computer
 - On completion of algorithm development the algorithm is ported to the DSP, or if necessary a separate embedded microcontroller

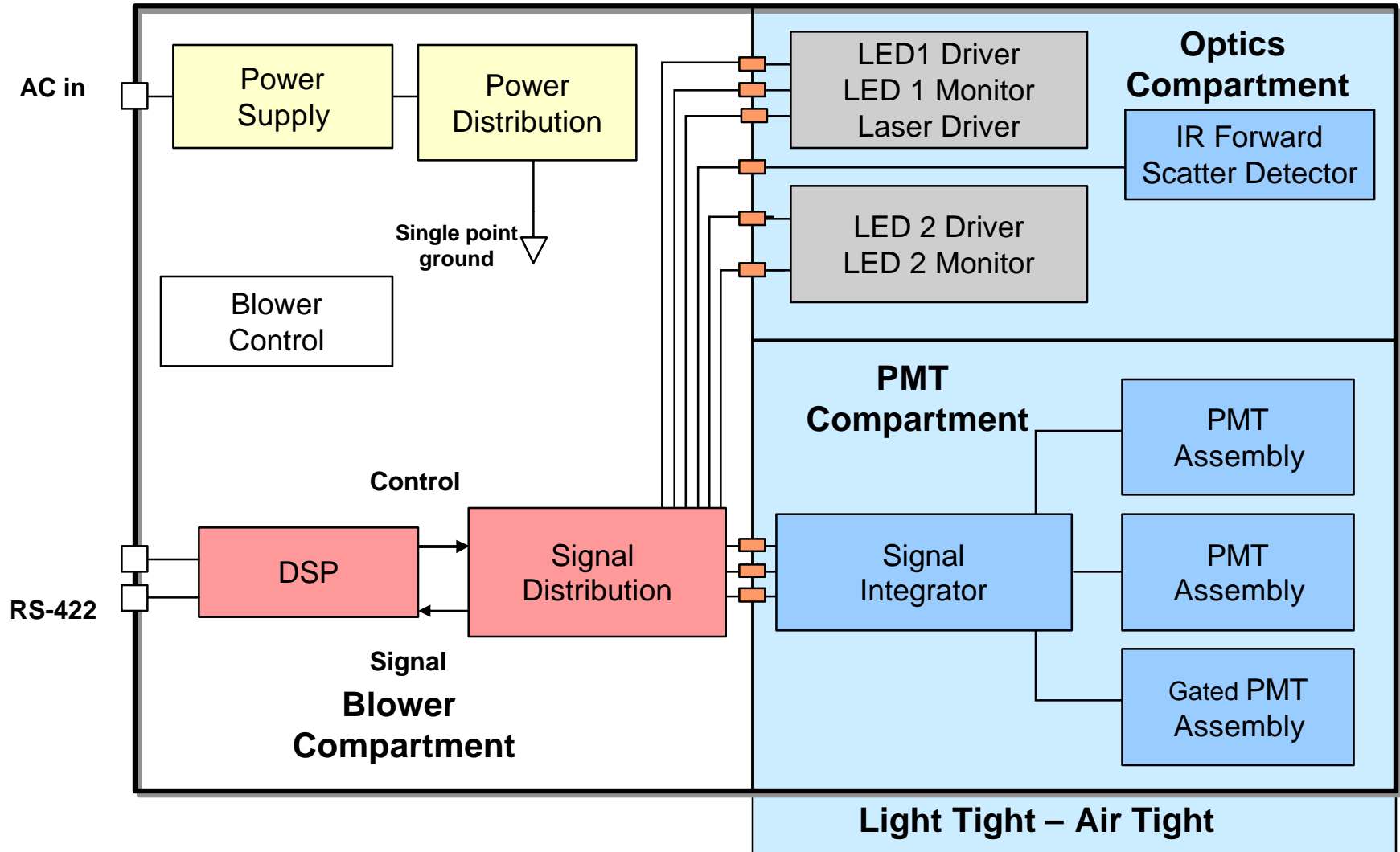


Electronics Block Diagram



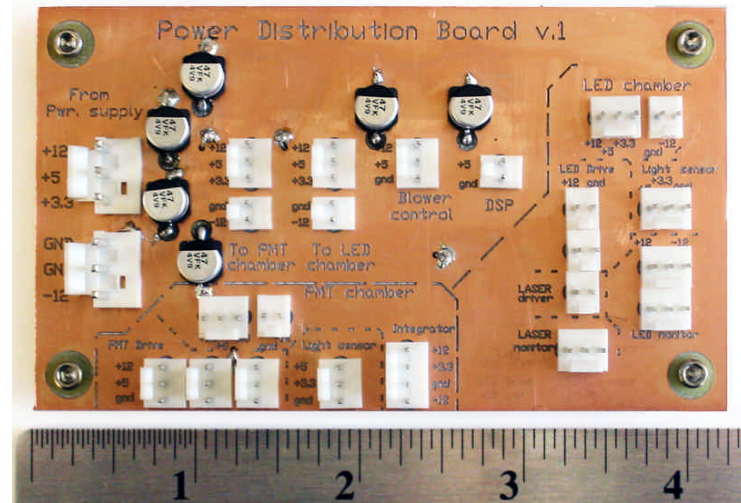


Electronics Board Configuration



[illegible]

- # Power Distribution Board



- **Output**
 - **Separate power for each of three compartments**
- **Filtering**
 - **Medium and high frequency noise filters**
 - **Ripple 20mv p-p, with fan running**
 - **Additional filtering on satellite boards**
- **Size**
 - **2.5" x 4.25" x .75"**



DSP/Microcontroller

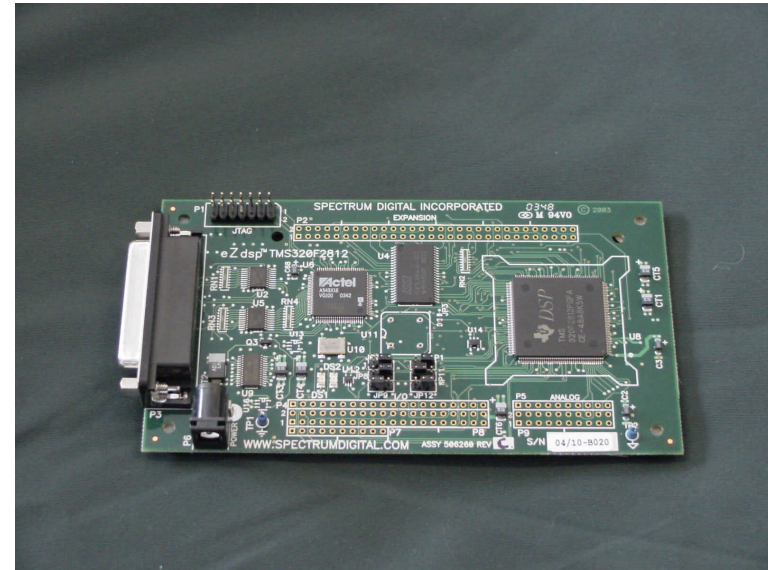
Texas Instruments TMS320F2812

- **Unifies microcontroller peripherals with DSP processing power**
 - 128K of flash and 18K of RAM
- **32-bit Harvard Architecture Processor with MAC**
 - Runs at 150 MHz for 6.67ns instruction time
- **On-board peripherals**
 - **Analog-to-Digital Converter**
 - 2 x 8 Channel 12.5 MSPS
 - 12-bit resolution
 - **Pulse width modulation outputs**
 - **56 general purpose digital ports**
 - 20 MHz modulation rate max
 - **Serial ports**
 - **Controller Area Network (CAN) Interface**
 - **Watchdog timer**
- **Real-time operating system**
- **Unit cost \$25 (bare chip)**



Digital Signal Processing Evaluation Board

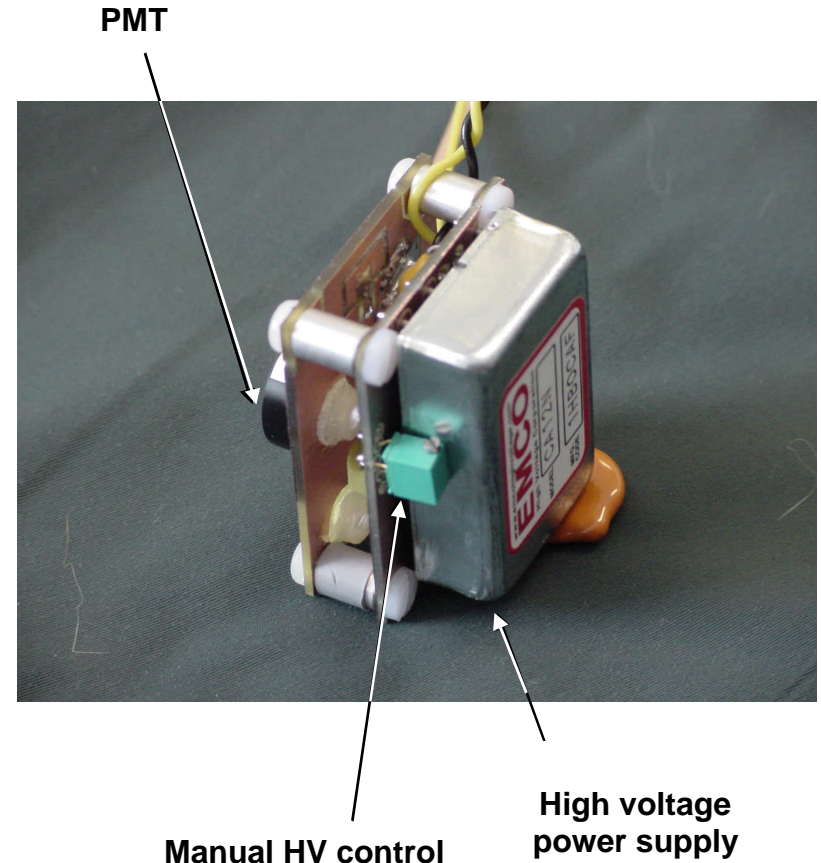
- **Spectrum Digital DSP evaluation board**
 - Interface headers
 - On board memory
 - JTAG ports
 - Cost ~\$450
- **Custom built distribution board**
 - Connects control and analog signals from DSP board to distributed analog boards





PMT Module

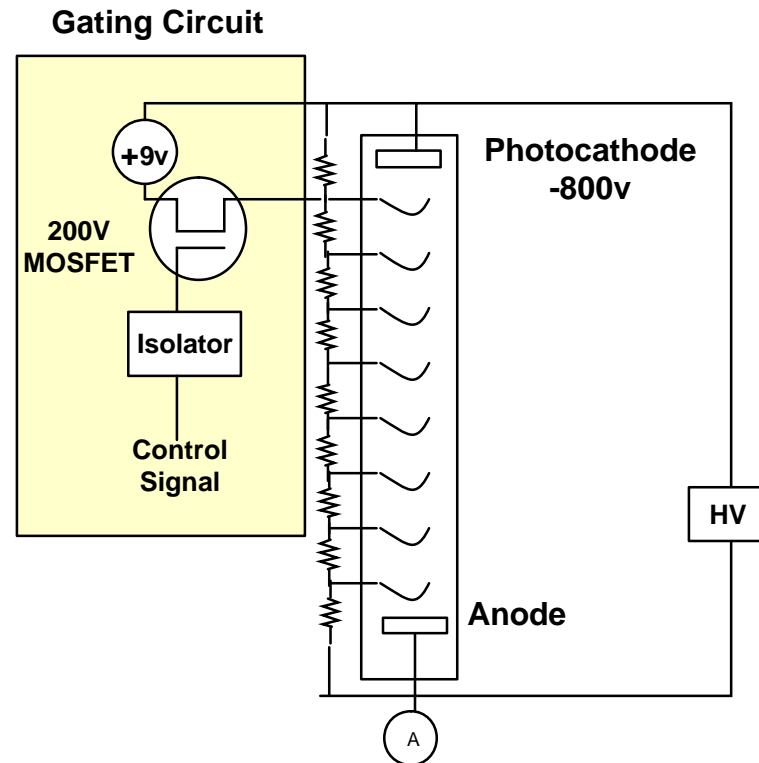
- **Photomultiplier Tube, Hamamatsu R7400-03**
 - Gain = 10^6
 - Dark current $<1\text{nA}$
 - Sensitivity = 62mA/W
 - Size = $1.75'' \times 0.75'' \times 0.5''$
 - Operating voltage = -800 VDC
- **Power Supply, EMCO CA12N**
 - 0 to -1250 VDC , 800mA max current
 - 12VDC input, $<220\text{ mA}$
 - Ripple $<0.0005\%$
 - Size = $1.75'' \times 0.75'' \times 0.5''$
- **Mounting**
 - Custom pair of boards designed for small form factor
 - Accurate alignment with light cone through mounting arrangement





Gated PMT Module

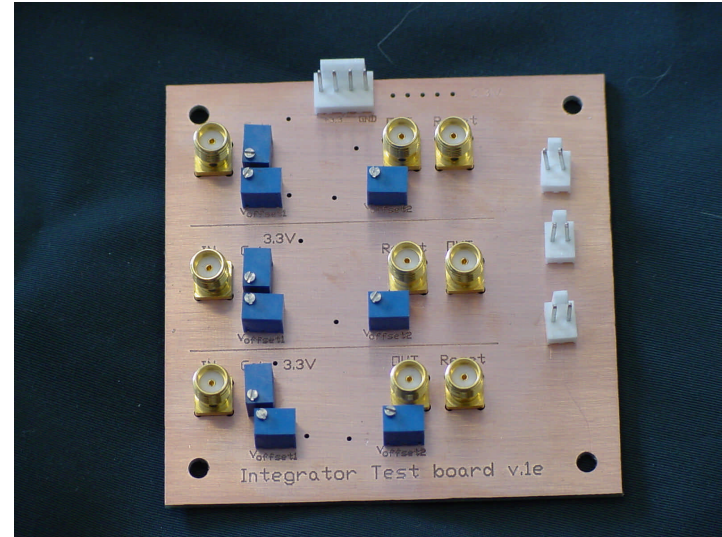
- **Gain reduction**
 - 100X
- **Switching speed**
 - < 4ms





PMT Signal Integrator

- 3 channel signal integration - one for each PMT
- Two gain settings for scatter and fluorescence measurements
- >3 log dynamic range in each channel
- Sample at beginning and end of integration cycle to reduce inaccuracies in reset
 - Measure when no signal present to determine offset current





LED Driver and Monitor

- **LED Driver**
 - Precision current source controlled by linear regulator and potentiometer
 - Switching rates of 5kHz up to 200mA
- **LED Monitor**
 - Transimpedance amplifier with adjustable gain
 - Particles < 1mm detectable



Data Flow at Signal Processor Output

- **Maximum particle rate = 100 particles / second**
 - Based on 1 ms pulse width
 - 10% duty cycle for UV LED
 - 10,000 ppl concentration
- **Particle characterization**
 - 10 descriptors
 - Timestamp, 4 bytes
Matlab format
- **Data format**
 - Fixed length record 20 bytes/particle
- **Signal rate**
 - Maximum ~2 KB/sec
175 MB/day
 - Typical (1000 ppl background) ~200 B/sec
17 MB/day



Summary

- **BAST electronics have been developed for support of a triggered, multi-band particle measurement instrument**
- **The electronics combines COTS and custom components**
- **No major issues have been uncovered in the development of the electronics**



Biological Agent Sensor Testbed Algorithm Development



Algorithm Development Goals

- **Develop real-time detection algorithm**
 - Detection of threat-like change in ambient aerosol
- **Operate in typical background conditions**
 - Good sensitivity
 - High probability of detection
 - Fast response time
 - Low false positive rate
- **Leverage previous experience with triggered point-detection bio-sensors and detection algorithms**



Data Flow to Algorithm

- **Particle Measurements**

- 280-nm elastic scattering
- 300 - 400 nm fluor. (280-nm ex.)
- 400 - 500 nm fluor. (280-nm ex.)
- 400 - 500 nm fluor. (340-nm ex.)
- IR scattering (820 nm)
- Particle transit time
- LED power
- Timestamp

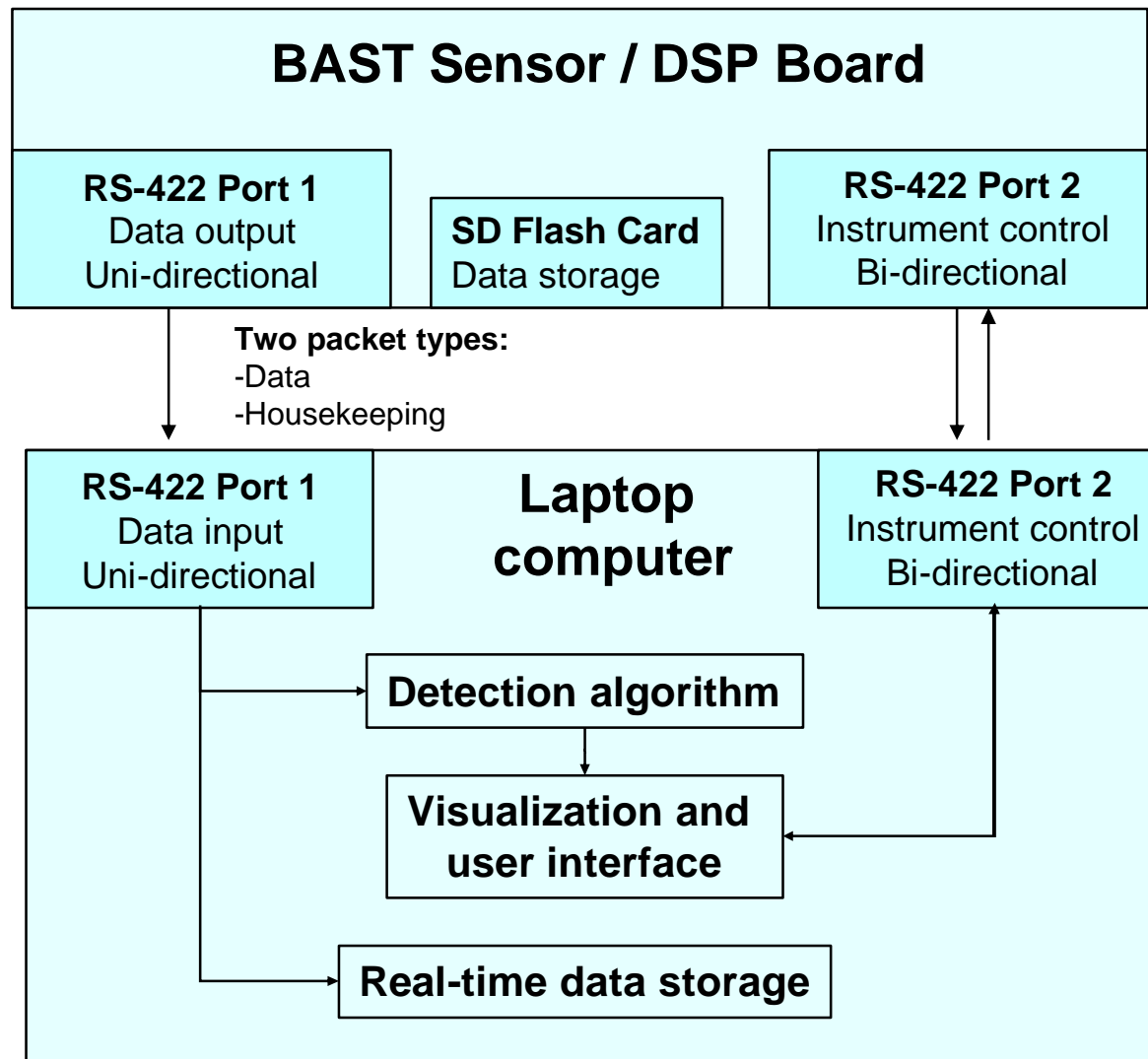
- **Particle triggered**
 - Raw particle signal
- **Periodic (1 Hz)**
 - Sensor noise
- **Particle signal = raw particle signal – sensor noise**

- **Housekeeping Measurements**

- Airflow
- LED operation time
- System voltages
- Etc...

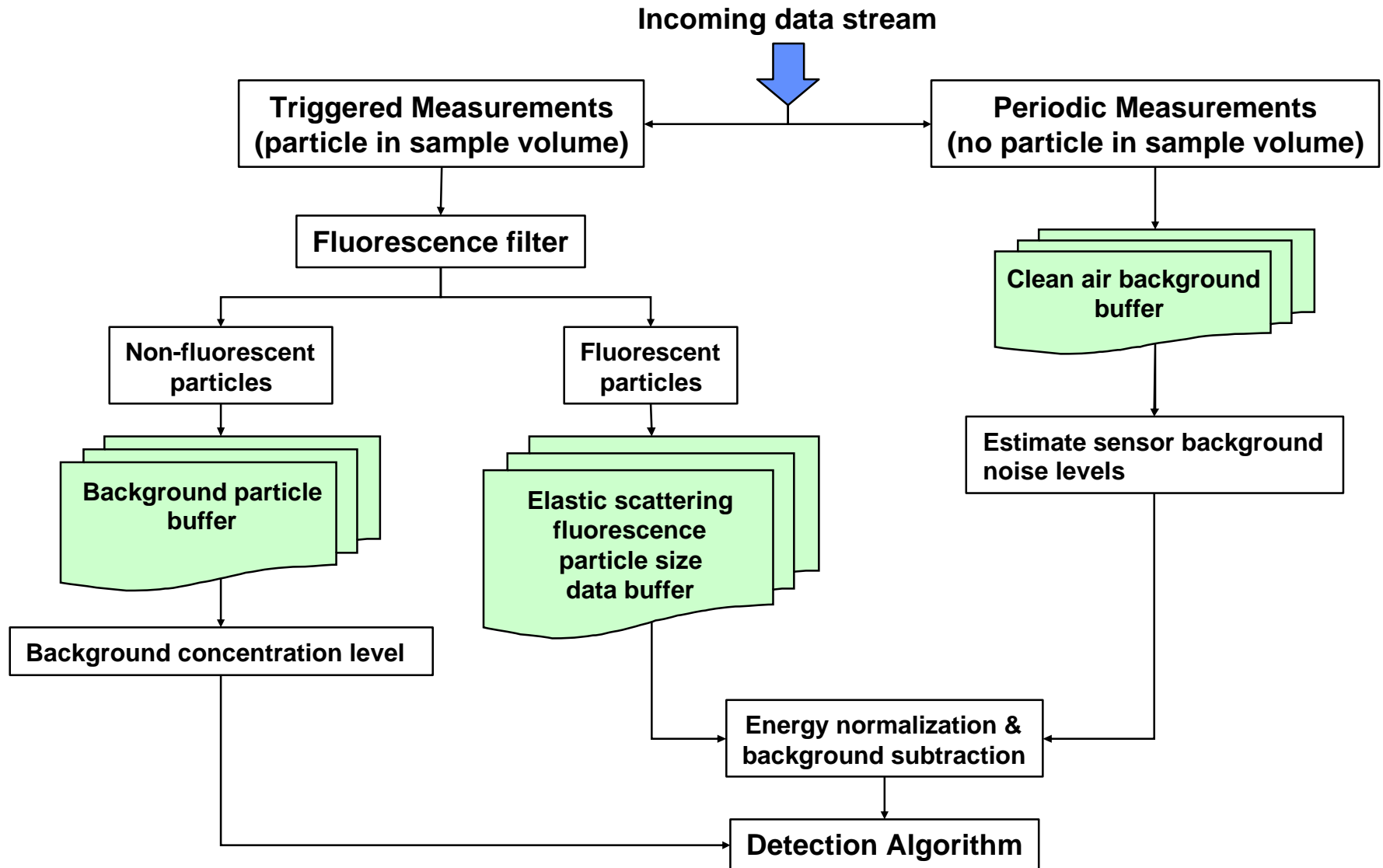


Data Collection and Instrument Control





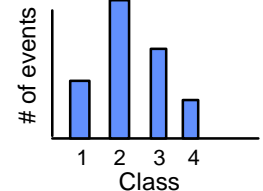
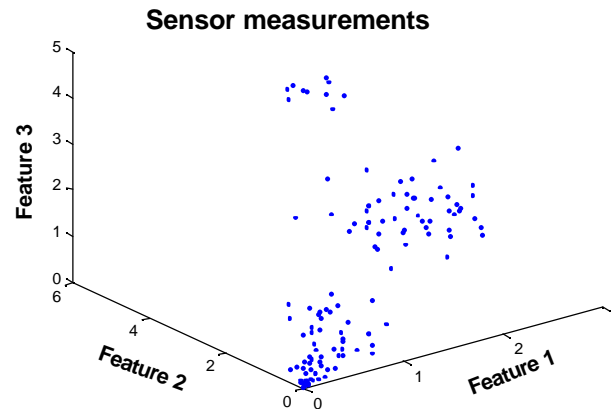
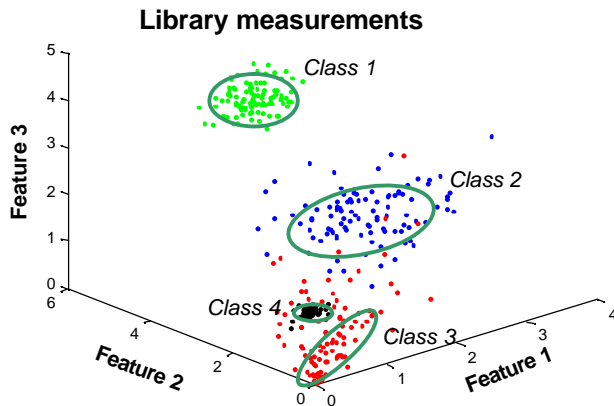
Algorithm Pre-processing





Algorithm Concept

- **Separate particles into broad classes based on position in feature space**
 - Directly measured features (e.g. elastic scattering, fluorescence, etc.)
 - Derived features (e.g. fluorescence/scattering ratio, spectral histogram, higher order statistics, etc.)
 - Regions defined by library of laboratory and field measurements
- **Track running histogram of occurrences for each class**
- **Positive detection if:**
 - Number of occurrence in any class exceeds predefined threshold count (100 particles)
- **Maximum response time one minute (minimum time about 10 seconds)**





High or Fluctuating Clutter Conditions

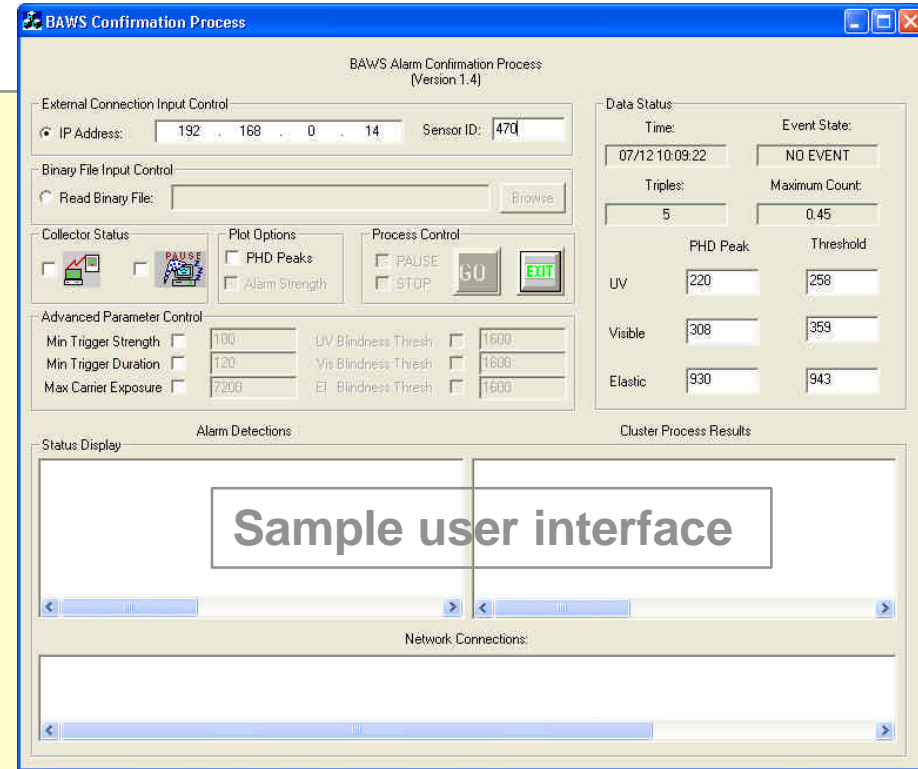
- **Ambient background conditions can vary dramatically with sensor location and operating environment**
- **Very high particle backgrounds can effectively “blind” the sensor to threat particles**
- **Track background concentration and correct measured threat particle concentration**
- **Several possible responses to this condition**
 - **Raise particle size threshold**
 - **Raise concentration threshold**
 - **Alert user to potential sensitivity reduction**



User Interface Development

Visualization and Analysis Tools

- **Development Interface**
 - Access to all relevant signals
 - Buffered analog test points
 - Digital outputs
 - Real Time Displays
- **Field Testing Interface**
 - Event data (standard detection algorithm output)
 - Real-time particle data
 - Summary statistics and time series
 - Diagnostics
- **End User Interface**
 - Trigger status
 - Sensor health status
 - Optional algorithm outputs over ethernet and/or serial port



- Verify incoming signals (“quick-look” data tools)
- Flexible visualization to support algorithm development
 - immediate feedback
 - modular



Summary

- **Developed an architecture for a real-time algorithm to detect potential threat aerosols**
- **Leverage prior experiences and lessons learned from previous bio-detection sensor systems and field measurements**
- **UV induced fluorescence has been shown be a good indicator of the presence of biological particles**
- **With five independent measurements we hope to increase discrimination between threat and non-threat particles**